

FORM PTO-1390 U.S. DEPARTMENT OF COMMERCE
PATENT AND TRADEMARK OFFICE (REV. 1094)

ATTORNEY'S DOCKET
NUMBER

C1043/7030

TRANSMITTAL LETTER TO THE UNITED STATES
DESIGNATED/ELECTED OFFICE (DO/EO/US)
CONCERNING A FILING UNDER 35 U.S.C. 371

U.S. APPLICATION NO. (If known, see 37 CFR 1.5)

09/743621

INTERNATIONAL APPLICATION NO.
PCT/GB99/02271

INTERNATIONAL FILING DATE
14 July 1999 (14.07.99)

PRIORITY DATE CLAIMED
14 July 1998 (14.07.98)

TITLE OF INVENTION
OPTICAL DEVICES

APPLICANT(S) FOR DO/EO/US

HO, Peter; TESSLER, Nir; FRIEND, Richard Henry

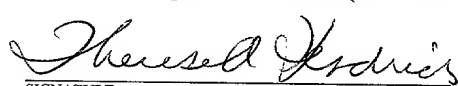
Applicant herewith submits to the United States Designated/Elected Office (DO/EO/US) the following items and other information:

1. ☒ This is a **FIRST** submission of items concerning a filing under 35 U.S.C. 371.
2. ☐ This is a **SECOND** or **SUBSEQUENT** submission of items concerning a filing under 35 U.S.C. 371.
3. ☒ This express request to begin national procedures (35 U.S.C. 371(f) at any time rather than delay examination until the expiration of the applicable time limit set in 35 U.S.C. 371(b) and PCT Articles 22 and 39(1).
4. ☒ A proper Demand for International Preliminary Examination was made by the 19th month from the earliest claimed priority date.
5. ☒ A copy of the International Application as filed (35 U.S.C. 371(c)(2)).
 - a. ☐ is transmitted herewith (required only if not transmitted by the International Bureau).
 - b. ☒ has been transmitted by the International Bureau.
 - c. ☐ is not required, as the application was filed in the United States Receiving Office (RO/US).
6. ☐ A translation of the International Application into English (35 U.S.C. 371(c)(2)).
7. ☒ Amendments to the claims of the International Application under PCT Article 19 (35 U.S.C. 371(c)(3)).
 - a. ☐ are transmitted herewith (required only if not transmitted by the International Bureau).
 - b. ☐ have been transmitted by the International Bureau.
 - c. ☐ have not been made; however, the time limit for making such amendments has NOT expired.
 - d. ☒ have not been made and will not be made.
8. ☐ A translation of the amendments to the claims under PCT Article 19 (35 U.S.C. 371(c)(3)).
9. ☐ An oath or declaration of the inventor(s) (35 U.S.C. 371(c)(4)).
10. ☐ A translation of the annexes to the International Preliminary Examination Report under PCT Article 36 (35 U.S.C. 371(C)(5)).

Items 11. To 16. Below concern document(s) or information included:

11. ☐ An Information Disclosure Statement under 37 CFR 1.97 and 1.98.
12. ☐ An assignment document for recording. A separate cover sheet in compliance with 37 CFR 3.28 and 3.31 is included.
13. ☒ A **FIRST** preliminary amendment (with amended claims).
 - ☐ A **SECOND** or **SUBSEQUENT** preliminary amendment.
14. ☐ A substitute specification (submitted as a first Preliminary Amendment).
15. ☐ A change of power of attorney and/or address letter.
16. ☒ Other items or information:
 - Mailed via Express Mailing Label No. EL711250388US
 - Published Application with Search Report
 - Post Card

Mailing Date: January 12, 2001
Express Mail Label No. EL711250388US

U.S. APPLICATION NO. (Re: PCT/PTO, see 37 CFR 1.492(a)(1)-(5))		INTERNATIONAL APPLICATION PCT/GB99/02271		ATTORNEY'S DOCKET NUMBER C1043/7030	
17. <input checked="" type="checkbox"/> The following fees are submitted:				CALCULATIONS PTO USE ONLY	
BASIC NATIONAL FEE (37 CFR 1.492(a)(1)-(5)): Search Report has been prepared by the EPO or JPO \$860.00 International preliminary examination fee paid to USPTO (37 CFR 1.482) \$690.00 No international preliminary examination fee paid to USPTO (37 CFR 1.482) but international search fee paid to USPTO (37 CFR 1.445(a)(2)).. \$760.00 Neither international preliminary examination fee (37 CFR 1.482) nor international search fee (37 CFR 1.445(a)(2)) paid to USPTO..... \$1000.00 International preliminary examination fee paid to USPTO (37 CFR 1.482) and all claims satisfied provisions of PCT Article 33(2)-(4) \$100.00					
ENTER APPROPRIATE BASIC FEE AMOUNT =				\$ 860.00	
Surcharge of \$130.00 for furnishing the oath or declaration later than ~ 20 X 30 months from the earliest claimed priority date (37 CFR 1.492(e)).				\$	
CLAIMS	NUMBER FILED	NUMBER EXTRA	RATE		
Total Claims	35 - 20 =	15	X \$18.00	\$ 270.00	
Independent Claims	3 - 3 =	0	X \$80.00	\$	
MULTIPLE DEPENDENT CLAIM(S) (if applicable)			+ \$260.00	\$	
TOTAL OF ABOVE CALCULATIONS =				\$1130.00	
Reduction by 1/2 for filing by small entity, if applicable. Verified Small Entity Statement must also be filed (Note 37 CFR 1.9, 1.27, 1.28).				\$	
SUBTOTAL =				\$1130.00	
Processing fee of \$130.00 for furnishing the English translation later than ~ 20 ~ 30 months from the earliest claimed priority date (37 CFR 1.492(f)).				\$	
TOTAL NATIONAL FEE =				\$1130.00	
Fee for recording the enclosed assignment (37 CFR 1.21(h)). The assignment must be accompanied by an appropriate coversheet (37 CFR 3.28, 3.31). \$40.00 per property +				\$	
TOTAL FEES ENCLOSED =				\$1130.00	
				Amount to be:	
				refunded	\$
				charged	\$
a. <input checked="" type="checkbox"/> A check in the amount of \$1130.00 to cover the above fees is enclosed. b. Please charge by Deposit Account No. _____ In the amount of \$ _____ To cover the above fees. A duplicate copy of this sheet is enclosed. c. <input checked="" type="checkbox"/> The commissioner is hereby authorized to charge any additional fees which may be required, or credit any overpayment to Deposit Account No. 23/2825. A duplicate of this sheet is enclosed.					
NOTE: Where an appropriate time limit under 37 CFR 1.494 or 1.495 has not been met, a petition to revive (37 CFR 1.137(a) or (b) must be filed and granted to restore the application to pending status.					
SEND ALL CORRESPONDENCE TO Therese A. Hendricks WOLF, GREENFIELD & SACKS, P.C. 600 Atlantic Avenue Boston, Massachusetts 02210					
SIGNATURE  NAME Therese A. Hendricks					
REGISTRATION NO 30,389					

ATTORNEY'S DOCKET NO: C1043/7030

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Applicant: Ho et al.
Serial No: --
Filed: Herewith
For: OPTICAL DEVICES

Assistant Commissioner for Patents
Washington, D.C. 20231

PRELIMINARY AMENDMENT

Please amend the above application as follows to conform the specification with U.S. practice.

In The Specification

- ◆ On page 1, before the first paragraph, please add the subheading -- Field of the Invention --
- ◆ On page 1, before the second paragraph, please add the subheading -- Background of the Invention --
- ◆ On page 4, before the last paragraph, please add the subheading -- Summary of the Invention --
- ◆ On page 10, before the second paragraph, please add the subheading -- Brief Description of the Drawings --;
- ◆ On page 11, before line 8, please add the subheading -- Detailed Description --

In the Abstract

Please add the following Abstract (also attached on a separate sheet at the end of this preliminary amendment):

with a solvent in which the nanoparticles are soluble to remove the said other material and form a solution of nanoparticles in the solvent. --

In The Claims

Please delete claims 36 and 37

Please amend the claims as follows:

- ◆ In claim 6 delete "any of claims 2 to 5" and substitute -- claim 1 --
- ◆ In claim 8, delete "any preceding" and after "claim" insert -- 1 --
- ◆ In claim 9, delete "any preceding" and after "claim" insert -- 1 --
- ◆ In claim 10, delete "any preceding" and after "claim" insert -- 1 --
- ◆ In claim 11, delete "any preceding" and after "claim" insert -- 1 --
- ◆ In claim 12, delete "any preceding" and after "claim" insert -- 1 --
- ◆ In claim 13, delete "any preceding" and after "claim" insert -- 1 --
- ◆ In claim 14, delete "any preceding" and after "claim" insert -- 1 --
- ◆ In claim 15, delete "any preceding" and after "claim" insert -- 1 --
- ◆ In claim 16, delete "any preceding" and after "claim" insert -- 1 --
- ◆ In claim 17, delete "any preceding" and after "claim" insert -- 1 --
- ◆ In claim 18, delete "any preceding" and after "claim" insert -- 1 --
- ◆ In claim 19, delete "any preceding" and after "claim" insert -- 1 --
- ◆ In claim 20, delete "any preceding" and after "claim" insert -- 1 --
- ◆ In claim 21, delete "any preceding" and after "claim" insert -- 1 --
- ◆ In claim 23, delete "or 22"
- ◆ In claim 24, delete "any of claims 1 to 20" and substitute -- claim 1 --
- ◆ In claim 26, delete "any of claims 1 to 20" and substitute -- claim 1 --
- ◆ In claim 27, delete "or 23"
- ◆ In claim 30, delete "any of claims 26 to 29" and substitute -- claim 26 --

- ◆ In claim 33, delete "any of claims 26 to 32" and substitute -- claim 26 --

Respectfully Submitted,



Therese A. Hendricks

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Date: January 12, 2001
Docket No. C1043/7030

C1043/7030 T294260

ABSTRACT

A method for preparing nanoparticles for use, from a mixture of nanoparticles with another material, the method comprising washing the mixture with a solvent in which the nanoparticles are soluble to remove the said other material and form a solution of nanoparticles in the solvent.

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T06040-T09E460

OPTICAL DEVICES

This invention relates to optical devices, especially devices comprising particles.

Nanoparticles are particles of very small size, typically less than 100nm across. The preparation of well-defined nanoparticles via colloid chemistry was demonstrated at least as early as the 1980s. A review of the current technology in this field is given in M P Pileni, *Langmuir*, 13, 1997, 3266-3279. There are three principal established routes for the formation of nanoparticles: a microemulsion route, a sol-gel route and a high temperature process used principally for semiconducting nanoparticles such as CdSe.

The synthetic principles of microemulsions have been widely described in the literature. For recent work see for example: F. J. Arriagada and K. Osseo-Asare, "Synthesis of Nanosized Silica in Aerosol OT Reverse Microemulsions," *Journal of Colloid and Interface Science* 170 (1995) pp. 8-17; V. Chhabra, V. Pillai, B. K. Mishra, A. Morrone and D. O. Shah, "Synthesis, Characterization, and Properties of Microemulsion-Mediated Nanophase TiO₂ Particles," *Langmuir* 11 (1995) pp. 3307-3311; H. Sakai, H. Kawahara, M. Shimazaki and M. Abe, "Preparation of Ultrafine Titanium Dioxide Particles Using Hydrolysis and Condensation Reactions in the Inner Aqueous Phase of Reversed Micelles: Effect of Alcohol Addition," *Langmuir* 14 (1998) pp. 2208-2212; J. Tanori and M. P. Pileni, "Control of the Shape of Copper Metallic Particles by Using a Colloidal System as Template," *Langmuir* 13 (1997) pp. 639-646. A microemulsion is a sufficiently thermodynamically stable solution of two normally immiscible liquids (for example, oil and water) consisting of nanosized droplets (or cores) of one phase in another "continuous" phase, stabilised by an interfacial film of a surfactant with or without a co-surfactant. Examples of surfactants include ionic ones such as Aerosol OT and cetyltrimethylammonium bromide, and non-ionic ones such as the polyoxyethylene ether and ester surfactants. Examples of co-surfactants include medium to long alkyl-chain alcohols such as 1-hexanol. Examples of oils include hydrocarbons such as cyclohexane and

isooctane. The surfactant and co-surfactant molecules reduce the interfacial tension so that stable dispersions can be formed.

Forming nanoparticles by the microemulsion route typically involves preparing a reaction mixture as a water-in-oil reverse micellar system using a ternary phase mixture containing high oil and surfactant contents, but low water content. This allows discrete but thermodynamically-stable nanometer-sized "water pools" or "water cores" to develop in the reaction mixture. In a typical water-in-oil microemulsion, the water cores are around 1 to 10nm in diameter. One reactant for the nanoparticle formation can be initially housed in these water cores. The second reactant can subsequently diffuse into and react inside these "nano-reactors" in the normal course of microemulsion dynamics. In this way, microemulsions provide a versatile route to the controlled synthesis of a wide array of oxide and non-oxide types of nanoparticle. In the water pools a metal salt can be reduced to the free metal, or metathesis reactions can be included, to obtain a controlled nucleation and growth of the desired nanoparticle material. The surfactant also acts as a coating to prevent unwanted flocculation (agglomeration) of the growing particles. Many of the fundamental principles governing such micellar chemistry, such as reaction rate and final growth size, are still largely unknown. Most experiments are done by trial-and-error and the data interpreted empirically.

Much of the work on nanoparticles has concentrated either: (i) on demonstrating that nanometer-sized particles have indeed been created (for instance by using transmission electron microscopy (TEM) or ultraviolet-visible (UV-Vis) spectroscopy); or (ii) on subsequently sintering the nanoparticles to prepare a sintered body. This work has involved relatively crude techniques for handling the nano-sized material. For aspect (i) there has generally been no need to isolate or further manipulate the nano-size material. For aspect (ii) the formed material has typically been recovered from the emulsion by bulk precipitation upon addition of a destabilising solvent, or by vacuum removal of the reaction solvent. The material is then sintered at high temperatures to obtain the desired nano-grained article after "burning off" of the

surfactant coating. Since the particles are to be sintered into a solid mass there is no need to counteract their tendency to aggregate.

Some work has been done on other uses for nanoparticles. S Carter, J C Scott and P J Brock, Appl. Phys. Lett, 71, 1997, 1145-1147 describe the use of polydispersed TiO_2 , SiO_2 and Al_2O_3 nanoparticles in the form of a blend in polymer LED devices with the aim of enhancing the forward emission of light generated in the LED and/or improving carrier injection and recombination. The route by which the nanoparticles are obtained is not described, but the particles are described as having relatively large sizes: 30 to 80nm, especially in comparison to the device thickness of 110nm. It appears from the presence of light scattering that the nanoparticle material suffers from agglomeration. Thus the nanoparticular nature of the material cannot be fully exploited.

In some other works, for instance V. L. Colvin, M. C. Schlamp & A. P. Alivisatos Nature 370, 6488 (1994) "Light-emitting-diodes made from cadmium diselenide nanocrystals and a semiconducting polymer", the use of CdSe nanocrystals as a form of transport layer (deposited neat either by spin-coating or electrostatic self-assembly) in a multilayered device with organic light-emitting polymers has also been described. Another reference relating to nanoparticle polymer composites is J. Schmitt, G. Decher, W. J. Dressick, S. L. Brandow, R. E. Geer, R. Shashidhar, and J. M. Calvert, "Metal nanoparticle/polymer superlattice films: fabrication and control of layer structure," *Adv. Mater.*, vol. 9, pp. 61, 1997.

Organic materials are used for a wide range of applications, including the formation of light emissive devices (see PCT/WO90/13148 and US 4,539,507, the contents of both of which are incorporated herein by reference). There is often a need to tune the properties of such an organic material. For example, in the manufacture of optoelectronic devices there is a need for control over various properties of the materials to be used, including conductivity (and/or mobility), refractive index, bandgap and morphology. Some examples of known techniques for tuning various properties are as follows:

1. Conductivity. This has been tuned by adding a chemical compound that acts as a donor or acceptor (namely an electronic dopant), see C. K. Chiang, C. R. Fincher, Y. W. Park, A. J. Heeger, H. Shirakawa, E. J. Louis, S. C. Gau, and A. G. MacDiarmid, "Electrical Conductivity in Doped Polyacetylene," *Phys. Rev. Lett.*, vol. 39, pp. 1098-1101, 1977.

2. Charge generation and photo-voltaic response. This has been tuned by blending two materials with appropriate electronic levels so that electrons prefer to reside on one and holes on the other. The blends have been either of two organic materials such as polymers (J. J. M. Halls, C. A. Walsh, N. C. Greenham, E. A. Marseglia, R. H. Friend, S. C. Moratti, and A. B. Holmes, *Nature* 376, 498 (1995), "Efficient photodiodes from interpenetrating polymer networks") or of organic material with a nano-particle (N. C. Greenham, X. G. Peng, and A. P. Alivisatos, "Charge separation and transport in conjugated-polymer/semiconductor- nanocrystal composites studied by photoluminescence quenching and photoconductivity," *Phys. Rev. B - Cond. Matt.*, vol. 54, pp. 17628-17637, 1996) to achieve exciton dissociation at the interface.

3. Band-gap and emission colour. This has been tuned by mixing organic compounds in the form of blends or co-polymers (see co-pending UK patent application number 9805476.0).

4. Scattering. Highly aggregated or very large size particles have been blended into polymers to take advantage of multiple internal light-scattering so as to increase the effective length the light travels within the polymer and hence, enhance amplified stimulated emission processes (F. Hide, B. J. Schwartz, M. A. Diazgarcia, and A. J. Heeger, "Laser-emission from solutions and films containing semiconducting polymer and titanium-dioxide nanocrystals," *Chem. Phys. Lett.*, vol. 256, pp. 424-430, 1996). Scattering might have also been used to enhance external efficiency of LEDs (S Carter, J C Scott and P J Brock, *Appl. Phys. Lett.* 71, 1997, 1145-1147).

According to a first aspect of the present invention there is provided a method for preparing nanoparticles for use, from a mixture of nanoparticles with another material, the method comprising washing the mixture with a solvent in which the nanoparticles are soluble to remove the said other material and form a solution of

nanoparticles in the solvent. The solvent is preferably one in which the said other material is not soluble. Alternatively, the solvent may be one in which the other material is soluble but the nanoparticles are not.

The method preferably comprises separating at least a first fraction of the nanoparticles from a mixture of the solvent and the said other material. These separated nanoparticles may then be used, for example in the applications described below. These nanoparticles are preferably only weakly bound (e.g. unaggregated or only weakly aggregated), and not strongly bound, so that they suitably exist in a disaggregated state. This can assist in subsequent processing steps, such as forming a substantially uniform dispersion of the nanoparticles in another material. The solvent is preferably one that is capable of holding the dissolved nanoparticles in a disaggregated state.

The method preferably includes a step of maintaining the pH of the solvent at a predetermined level. This suitably maintains a charge on the nanoparticles. An acid or a base and/or a suitable buffer may be added to the solvent to maintain the desired pH.

The method preferably includes dialysis through a suitable membrane to remove soluble low molecular weight material (for example surfactant molecules) from the nanoparticle solution. Continuous or intermittent sonification could be performed during dialysis.

The separation may be performed by filtration and/or dialysis and/or centrifugation. Preferably the separation step or another step of the method also allows for the separation of the said first fraction of the nanoparticles from another fraction of the nanoparticles. For example, the nanoparticles of the said other fraction may be a set of particles that are relatively small in comparison to the nanoparticles of the first fraction. Thus, the separation step may also serve to narrow the size distribution of the retained nanoparticles.

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The method preferably comprises surface modifying the nanoparticles. This may suitably be achieved by adsorbing a material to the surface of the particles. The material may be added as surface modifying agent to the solution of nanoparticles. The material may, for example, be a silylating agent or a dye or a chemical functional material. The material may promote specific interactions with other materials such as polymers. Alternatively, or additionally, the nanoparticles may already have a surface coating. This may be a coating of a surfactant.

The nanoparticles may be of metallic, semiconducting or insulating material. Examples of suitable materials include inorganic oxides such as SiO_2 , TiO_2 , Al_2O_3 or ZrO_2 , or ternary or other binary inorganic materials such as BaSO_4 , YbF_3 , ZnS or other organic materials, especially polymer materials, such as PTFE, polymethylmethacrylate (PMMA) or polystyrene (PS). The nanoparticles are preferably

light transmissive and most preferably optically transparent. Therefore, the material of which the nanoparticles are formed is preferably a wide optical bandgap material.

The nanoparticles may have been formed by any suitable route. Examples include the microemulsion route and the sol-gel route.

A further step in the processing of the nanoparticles is preferably to incorporate them into a body of material. To achieve this the material of the body, or a precursor of it, is preferably added to the solution of nanoparticles. A uniform (or substantially uniform) non-aggregated (or substantially non-aggregated) dispersion of the nanoparticles in the final body is achieved by ensuring that they are held in a substantially disaggregated state until fixed in place in the body, e.g. by removal of the solvent by a step such as evaporation. It is therefore preferable that the material of the body (or its precursor) is soluble in the solvent in which the nanoparticles are dissolved, and does not have undesirable interactions with the nanoparticles that may lead to severe aggregation or phase separation.

The material of the body could be (but need not be) an organic material. Examples are polymers, oligomers and materials of small organic molecules. If the material is a polymer material it may be a conjugated polymer such as poly(p-phenylenevinylene) (PPV). Alternatives include poly(2-methoxy-5-(2'-ethyl)hexyloxyphenylene-vinylene) ("MEH-PPV"), a PPV-derivative (e.g. a di-alkoxy or di-alkyl derivative), a polyfluorene and/or a co-polymer incorporating polyfluorene segments, PPVs and/or related co-polymers, poly(2,7-(9,9-di-n-octylfluorene)-(1,4-phenylene-((4-secbutylphenyl)imino)-1,4-phenylene)) ("TFB"), poly(2,7-(9,9-di-n-octylfluorene) - (1,4-phenylene-((4-methylphenyl)imino)-1,4-phenylene-((4 - methylphenyl)imino) - 1,4-phenylene)) ("PFM"), poly(2,7 - (9,9 - di-n-octylfluorene) - (1,4-phenylene-((4-methoxyphenyl)imino)-1,4-phenylene- ((4-methoxyphenyl)imino)-1,4-phenylene)) ("PFMO"), F8 or F8BT. Alternative materials include organic molecular materials such as Alq3. The material is suitably light-transmissive and/or light-emissive.

The presence of the nanoparticles (including optionally any material attached to the surface of the nanoparticles) in the material of the body preferably influences at least one material property of the material of the body. This could be an optical property such as refractive index or an electrical property such as conductivity. Thus the nanoparticles could be dispersed in the material of the body to tailor its refractive index - either increasing or decreasing it depending on the relative refractive indices of the material of the body and the nanoparticles. The presence of the nanoparticles, and the interaction between the nanoparticles and the polymer (including optionally any material attached to the surface of the nanoparticles) in the material of the body could also influence the morphology of the material of the body, for instance by inhibiting crystallisation.

The volume fraction of the nanoparticles in the body is preferably greater than 1, 5 or 10 volume %. The volume fraction of the nanoparticles in the body is preferably less than 50, 30 or 30 volume %. The density of the nanoparticle distribution in the body is preferably greater than 10^{-17} and/or less than $10^{-19}/\text{cm}^2$.

It is preferred that the nanoparticles exist in the body in a disaggregated state. This suitably promotes uniformity of the properties in the body. Furthermore, in some circumstances aggregates of the particles could scatter incident light, and it is preferred that the particles are of a sufficiently small size and disaggregated nature that they substantially do not scatter incident light.

The body could be a layer of a device such as an electronic and/or optical device. Preferred non-limiting examples of such devices are as follows:

1. A device comprising a stack of layers defining a light-reflective structure, with at least one of the layers comprising a dispersion of nanoparticles as described above. Preferably alternating layers of the device comprise a dispersion of nanoparticles as described above. This device is suitably a distributed Bragg reflector.
2. A light emissive device in which a light emissive layer or a layer adjacent to the light emissive layer comprises a dispersion of nanoparticles as described above. In this case the nanoparticles may carry a fluorescent dye (suitably as a surface layer).

This dye can suitably be stimulated to fluoresce by energy transfer or light emission from the emissive layer; thus the dye can act to modify the colour of light emission from the emissive layer. The device may also comprise a waveguide structure defined by a relatively high refractive index layer located between two relatively low refractive index layers. One of those three layers preferably comprises a dispersion of nanoparticles as described above which suitably modifies its refractive index to help define the waveguide structure. The waveguide is preferably located outside and/or separately from and/or independent of the light-emissive region of the device (the energy level profile of the device may be arranged to encourage light emission other than in the waveguide). This can permit independent tuning of the material properties of the emissive layer and the waveguide layer. The device may have a pair of mirrors located on either side of it - either mirrors of the type described for device 1 above or mirrors of another type such as cleavage surfaces. These mirrors may define a microcavity which can spectrally redistribute light generated by the device. The device may be a laser, for instance a microcavity or waveguide laser. A mirror could be provided by a DBR grating superimposed on the waveguide structure (on a substrate or any subsequent layer).

Other aspects of the present invention include any or all of the articles described above. For example, according to a second aspect of the present invention there is provided an optical and/or electronic device including any of the features described above. According to a third aspect of the invention there is provided a method for forming an optical and/or electronic device including any of the features described above. According to a fourth aspect of the invention there is provided a solution of nanoparticles in other than a strongly bound state, suitably including any of the features described above. According to a fifth aspect of the invention there is provided a solution of a polymer material (or a polymer precursor material) and nanoparticles in other than a strongly bound state. According to a sixth aspect of the invention there is provided an organic material containing a substantially uniform dispersion of nanoparticles; preferably the organic material is a semiconductive and/or a polymer material. According to a seventh aspect of the invention there is provided a method for tailoring at least one property of an organic material, the

method comprising forming a substantially uniform dispersion of nanoparticles in the material.

The present invention will now be described by way of example with reference to the accompanying drawings, in which:

figure 1 shows the results of energy dispersive X-ray spectrometry of TiO_2 nanoparticles;

figure 2 shows the results of Raman spectroscopy of TiO_2 nanoparticles;

figure 3 shows the results of attenuated total reflection infrared spectroscopy of TiO_2 nanoparticles;

figure 4 shows the UV-Vis spectrum of TiO_2 nanoparticles;

figure 5 shows the results of transmission electron microscopy of TiO_2 nanoparticles deposited as a film;

figure 6 shows an electron diffraction pattern for TiO_2 nanoparticles;

figure 7 shows the results of energy dispersive X-ray spectrometry of SiO_2 nanoparticles;

figure 8 shows the results of Raman spectroscopy of SiO_2 nanoparticles;

figure 9 shows the results of attenuated total reflection infrared spectroscopy of SiO_2 nanoparticles;

figure 10 shows the UV-Vis spectrum of SiO_2 nanoparticles;

figure 11 shows the results of transmission electron microscopy of SiO_2 nanoparticles deposited as a film;

figure 12 shows an electron diffraction pattern for SiO_2 nanoparticles;

figure 13 plots refractive index measurements for PPV: TiO_2 thin films;

figure 14 illustrates the structure of a distributed Bragg reflector;

figures 15 and 16 each show reflection spectra for two distributed Bragg reflectors showing that the reflection peak can be tuned;

figure 17a shows the structure of an edge emitting organic light emitting device;

figure 17b shows the refractive indices (n) and energy levels (E) through the device of figure 17a;

figure 18 shows difference UV-Vis spectra for PPV containing dye-loaded nanoparticles;

figure 19a shows the result of atomic force microscopy of a PPV film;

figure 19b shows the result of atomic force microscopy of a PPV:TiO₂ film with 30% vol. TiO₂; and

figure 20 shows UV-Vis spectra for PPV:TiO₂ films with differing amounts of TiO₂ nanoparticles.

Processes for the formation of nanoparticles, together with some exemplary applications for those nanoparticles (or for other suitable nanoparticles), will now be described. The processes allow for the formation of nanoparticles in a sufficiently non-aggregated state that they can be dispersed relatively evenly into a matrix such as a polymer body. In that state the nanoparticles can be employed to tailor the properties of the matrix in desired ways.

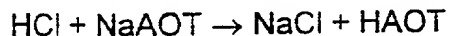
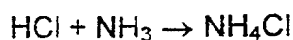
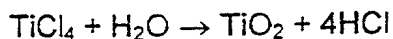
Processes for the synthesis and isolation of TiO₂ and SiO₂ nanoparticles will be described. These processes can readily be extended to the synthesis and isolation of nanoparticles of other materials, including metallic and semiconductive materials.

Synthesis of TiO₂ Nanoparticles

As a first step, a microemulsion of H₂O+NH₃/AOT/cyclohexane was prepared. To prepare the microemulsion 17.2g of finely divided sodium dioctylsulfosuccinate (AOT) (39mmol, Aldrich) was dried at 120°C in a vacuum oven for 2h to remove adsorbed moisture (3% weight loss), and then added to 200ml of HPLC grade cyclohexane (Aldrich) in a three-necked round bottom flask quipped with a magnetic stirrer to give a 0.2M AOT solution. Then 3.8ml of concentrated aqueous NH₃ (35w/v%, Aldrich) was added to this solution to give a H₂O:NH₃:AOT molecular ratio of 1.8:3.2:1.0 and the solution was cooled to 5°C under N₂ flow in an ice-bath to give an optically clear single-phase microemulsion. The total molar ratio of H₂O+NH₃:AOT (5.0:1.0) fixed the radius of water cores in the emulsion at around 25Å (F. J. Arriagada and K.

Osseo-Asare, "Synthesis of nanosize silica in Aerosol OT reverse microemulsions", Journal of Colloid and Interface Science 170 (1995), pp. 8-17).

To prepare a TiO_2 precursor solution, 2.0ml TiCl_4 (18mmol, Aldrich) was added by syringe into 10ml HPLC grade cyclohexane (previously dried for 18h over 3Å molecular sieves, Aldrich) in a vial equipped with a rubber septum. This TiO_2 precursor solution was added in two portions to the AOT solution with vigorous stirring. Rapid evolution of HCl gas was observed together with the formation of a cloudy suspension of NH_4Cl in the reaction medium, according to the following equations:



After half an hour, the bath temperature was raised to and then maintained at 80°C for 3h to complete the reaction, and finally cooled to room temperature overnight.

The amount of H_2O in the reaction medium (120mmol) was well in excess of the amount needed to hydrolyse the TiCl_4 . However, the amount of NH_3 was not sufficient to neutralise all the HCl produced, so the reaction medium became acidic and a positive surface charge developed on the TiO_2 nanoparticles. (The iso-electric point of bulk TiO_2 is about pH 4 to 5).

Isolation of TiO_2 Nanoparticles

The cyclohexane solvent was slowly evaporated away to give a cloudy, viscous gel containing the TiO_2 and NH_4Cl by-product. 20ml HPLC-grade CHCl_3 (Aldrich) was added to this gel to precipitate out the NH_4Cl and NaCl which could then be readily separated from the clear supernatant by centrifugation. Electron beam elemental analysis by energy dispersive x-ray spectrometry (EDX) of the CHCl_3 supernatant gave no detectable Cl (the height of the Cl Kα peak was less than 1% of the height of the Ti Kα peak), indicating it was essentially free from contamination by the chlorides.

The CHCl_3 supernatant was then concentrated *in vacuo* to give a viscous gel which was then taken up in 25ml MeOH to give the crude TiO_2 -containing liquor.

To purify the TiO_2 further from excess AOT, 10ml of the crude liquor was placed in a 6-8k MWCO cellulose dialysis tubing (Spectra/Por®) and dialysed against twice-daily refreshed 200ml quantities of HPLC grade MeOH each containing 0.3ml concentrated aqueous HCl. The Ti, S and Cl content of both the dialysate and retentate were monitored daily by EDX, and the dialysis was stopped when the Ti and S signals in the dialysate fell below that of Cl (which was treated as indicating a background level). This dialysis, in addition to removing the excess AOT contamination, also improved the size distribution of the particles by removing the smallest particles, which could readily pass through the pores of the dialysis tubing. In this way, an optically clear dispersion of about 0.9w/v% TiO_2 in a MeOH solution of pH=2 was obtained. The TiO_2 nanoparticles at this point were para-crystalline and coated by a monolayer of surfactant derivative.

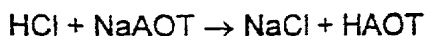
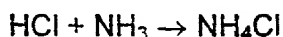
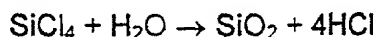
Characterisation of TiO_2 Nanoparticles

Electron probe energy dispersive x-ray spectrometry (see figure 1) shows the final product to be pure TiO_2 with a surface coating of AOT and Cl. It is believed that the AOT (which is the origin of the S signal) is strongly absorbed onto the TiO_2 surface while Cl is believed to have existed in the form of Cl^- acting as a weakly-bound counterion for the positively-charged TiO_2 surface. This coating appears to be stable against further dialysis. Raman spectroscopy (see figure 2) confirms the structure as being that of TiO_2 , with broad vibration peaks at 170cm^{-1} , 440cm^{-1} and 610cm^{-1} . Attenuated total reflection Fourier-transformed infrared spectroscopy (see figure 3) provides further confirmation of structure. Ultraviolet-visible spectrometry (see figure 4) shows the product to be non-scattering in the visible, and provides evidence for the highly dispersed (non-aggregated) nature of the TiO_2 nanoparticles achieved via the above described synthesis and isolation. Transmission electron microscopy (see figure 5) shows that a particle size of 2-8nm has been achieved, while electron beam diffraction (see figure 6) gives the expected d-spacings for TiO_2 .

Synthesis of SiO₂ nanoparticles

An H₂O+NH₃/AOT/cyclohexane phase was prepared as described above.

To prepare a SiO₂ precursor solution, 2.1ml SiCl₄ (18mmol, Aldrich) was added by syringe into 10ml HPLC grade cyclohexane (previously dried for 18h over 3Å molecular sieves, Aldrich) in a vial equipped with a rubber septum. This SiO₂ precursor solution was added in two portions to the AOT solution with vigorous stirring. Rapid evolution of HCl gas was observed together with the formation of a cloudy suspension of NH₄Cl in the reaction medium, according to the following equations:



After half an hour, the bath temperature was raised to and then maintained at 80°C for 3h to complete the reaction, and then cooled to room temperature overnight.

The amount of H₂O in the reaction medium (120mmol) was well in excess of the amount needed to hydrolyse the SiCl₄.

Isolation of SiO₂ nanoparticles

The cyclohexane solvent was slowly evaporated away to give a cloudy, viscous gel containing the SiO₂ and NH₄Cl (together with NaCl) by-product. Then 20ml HPLC-grade CHCl₃ (Aldrich) was added to this gel to precipitate out the SiO₂ together with NH₄Cl and NaCl by-product as a slurry which could then be readily separated from the clear supernatant which contained mainly excess AOT by centrifugation. EDX of the CHCl₃ supernatant gave no detectable Si (the height of the Si Kα peak was less than 1% of the height of the S Kα peak), indicating that it was essentially free from SiO₂. The CHCl₃ supernatant was then removed, and the slurry washed (with sonication) in several 25ml lots of HPLC-grade MeOH to remove as much NH₄Cl and

NaCl by-product as possible. After each washing the amount of SiO_2 in solution was checked. After three washings the centrifugate containing the SiO_2 was then taken up in 30ml MeOH, to give the crude SiO_2 as a translucent gel.

To purify the crude SiO_2 further from the inorganic chlorides, 10ml of the gel was placed in a 6-8k MWCO cellulose dialysis tubing (Spectra/Por®) and dialysed against two change of 200mL HPLC grade MeOH containing 0.3ml concentrated aqueous HCl. The retentate was then sonicated for 30min to break up the weak SiO_2 aggregates and then dialysis continued with twice-daily refreshed MeOH+HCl. The appearance of the retentate became clear after a while. The Si, S and Cl content of the dialysate was monitored daily, and the dialysis was stopped when the Cl signal fell to and stabilised at about 1.7 times that of S (from the AOT) which acted as a background. Sonication could be performed during dialysis. This dialysis, in addition to removing the chloride and AOT contamination, also improved the size distribution of the particles by cutting off the smallest particles, which could readily pass through the pores of the dialysis tubing. The dialysis took three to five days. A further dialysis step, with a larger molecular weight cut-off tube, could be used to separate off the largest particles and thus give an even tighter size distribution.

In this way, an almost optically clear dispersion of SiO_2 of about 1.1w/v% in a MeOH solution of pH=2 was obtained.

Characterisation of SiO_2 nanoparticles

Electron probe energy dispersive x-ray spectrometry (see figure 7) shows the final product to be pure SiO_2 with essentially no surface coating. Raman spectroscopy (see figure 8) confirms the structure as being that of SiO_2 with broad vibration peaks at 490cm^{-1} , and 830cm^{-1} . Attenuated total reflection Fourier-transformed infrared spectroscopy (see figure 9) provides further confirmation of structure. Ultraviolet-visible spectrometry (see figure 10) shows the product to be essentially non-scattering in the visible, providing evidence for the highly-dispersed (non-aggregated) nature of the SiO_2 nanoparticles achieved in the described synthesis and isolation.

Transmission electron microscopy (see figure 11) shows that a particle size range of 2-8nm was been achieved, while electron beam diffraction (see figure 12) gives the expected d-spacings for SiO₂.

General Nanoparticle Considerations

The nanoparticle synthesis process can be suited to formation of nanoparticles of a desired type by selection of the materials involved in the process, for instance the precursor materials and surfactant, and optimisation of chemical parameters such as oil composition, temperature, water-surfactant ratio, surfactant-oil ratio, co-surfactant composition and reactant ratio. Nucleation and growth of the particles could be monitored by TEM and/or UV-Vis spectroscopy to ensure suitable reaction rates and size distribution are achieved.

It is preferable that the steps be performed in a controlled atmosphere to minimise undesirable reaction in the nanoparticle solution. The temperature of the nanoparticle formation step could be fixed to favour the desired balance between nucleation and growth rates to give a preferred particle size distribution. It will be noted that the emulsion conditions described above have been chosen to yield especially small water cores.

The synthesis processes described above make use of the microemulsion route, which has been found generally to offer good control of particle nucleation, growth, aggregation, size distribution, and surface capping that could be judiciously exploited to tailor specific material and processing properties. As an alternative the nanoparticles could be synthesised by other routes, such as the sol-gel route based on direct hydrolysis in water or alcohols, (see for example T. Moritz, J. Reiss, K. Diesner, D. Su, and A. Chemseddine," Journal of Physical Chemistry B 101 (1997) pp. 8052-8053) and then isolated in a non-strongly aggregated state generally as described above.

Applications for the Nanoparticles

Some non-limiting examples of applications of the nanoparticles as prepared above will now be described. Nanoparticles prepared by other routes could be used in these applications, provided they were of the appropriate size and composition and had a level of disaggregation sufficient to allow them to be dispersed relatively uniformly in the polymer matrices.

Two general modes have been identified by which the nanoparticles can provide an effect in a host matrix material.

1. Non-interfering mode. The particles may be selected and/or treated so that they do not significantly alter the intrinsic properties of the host material. For most applications it would be preferred that such particles are of a size below around 20nm (to allow them to be easily dispersed uniformly in the host) and above 5nm (suitably to avoid them acting like molecules and possibly interfering with the host structure - for example by chain separation in a polymer matrix).

2. Interfering mode. The particles may be selected and/or treated so that they do interact significantly with the host material to affect its intrinsic properties. This may occur if:

- a. The particles themselves or especially their attached surface functions are of a material that interacts with the host material, through chemical or physical interactions. The particles may carry an additive agent that interacts with the host material, for example to change the chemical properties of the host (e.g. electrical/optical doping) and/or its physical properties (e.g. polarisation or alignment or energy transfer). The additive agent may be bound to the surface of the particles or at least partially included in the particles.
- b. The particles are sufficiently small and numerous that they significantly perturb the arrangement of the molecules in the body - for instance to affect the morphological structure of a polymer host (e.g. by perturbing the chain trajectories). Such particles could also change the environment of molecules of the host material (e.g. polymer chains) thus affecting many of its properties (band-gap, electron affinity, exciton binding, radiative lifetime etc.).

Example 1: Tuning of Polymer Film Refractive Index

Four samples of PPV containing varying amounts of TiO_2 (referred to herein as "PPV: TiO_2 ") were fabricated by blending the appropriate volume of the 1.4w/v% precursor PPV-MeOH solution (Cambridge Display Technology) with a 1.8w/v% TiO_2 -MeOH solution to give the same final concentration of precursor PPV but different concentrations of TiO_2 . The size range of the TiO_2 particles was approximately 2-8nm. A nominally 1000Å film was obtained by spin-coating on to a glass substrate and then thermally-converted at 180°C under dynamic vacuum (less than 10^{-5} mbar) for 8h. The actual TiO_2 content in the film was measured by a combination of electron probe x-ray spectrometry (for TiO_2 content) and visible spectrometry (for PPV content) and the film thickness was measured by surface profilometer.

Film	PPV:MeOH:TiO ₂ Vol. Ratio	Measured TiO ₂ Vol% in Final Film	Measured Refractive Index at 633nm
A	3:4:0	0	2.60
B	3:1:3	15	2.37
C	3:2:2	30	1.94
D	3:0:4	50	1.77

The refractive indices of the films over the whole visible window were then measured by spectroscopic ellipsometry (see figure 13). Transmission spectrometry shows the resultant films to be highly non-scattering (scatter at 633nm was estimated at less

than 2%). These results demonstrate that high optical-quality organic films can be obtained using highly-dispersed nanoparticles as a refractive index modifier.

Similar results were obtained for samples of PPV blended with SiO₂ (PPV:SiO₂). In the films detailed in the following table, the precursor PPV and SiO₂ concentrations were both 1.1w/v%. Nominally 1000Å films were obtained by repeated spin-coating and conversion on glass substrates and then thermally-annealing at 160°C under dynamic vacuum (less than 10⁻⁵ mbar) for 8h. The other experimental procedures were similar to those given earlier.

Film	PPV: MeOH: SiO ₂ Vol. Ratio	Measured SiO ₂ Vol% in Final Film	Measured Refractive Index at 633nm
E	3:16.5:0	0	2.39
F	3:15.5:1	6	2.18
G	3:14.5:2	10	2.10
H	5:12:4	19	2.05

To further check the effect on refractive index, films of similar thickness were placed between two silver mirrors to form a Fabry-Perot cavity. Since the resonance frequency of the cavity is a function of the optical length it serves as a measure for the effective refractive index.

The precursor PPV polycation and the negatively-charged SiO₂ nanoparticles interact to give a precipitate when mixed together. However, concentration conditions could be selected to overcome this problem (with low PPV and SiO₂ concentrations relative to the MeOH), and any slight precipitation could be removed by centrifugation. The optically-clear centrifugate could then be used to fabricate films with almost no scattering.

Photoluminescence measurements indicate that high photoluminescence efficiency is retained in the PPV:SiO₂ films. in contrast to PPV:TiO₂ films since, unlike TiO₂ particles, SiO₂ particles do not quench the PPV excited states. PPV:SiO₂ films could

therefore be preferred for use as emissive layers of electroluminescent devices. (Alternatively, quenching nanoparticles such as TiO_2 could be treated to carry a surface isolation or spacing layer of a material such as alkyl chains or SiO_2). The basic structure of such devices is well known and generally comprises a light-emissive organic layer, for instance a film of PPV, sandwiched between two electrodes. One of the electrodes (the cathode) injects negative charge carriers (electrons) and the other electrode (the anode) injects positive charge carriers (holes). The electrons and holes recombine in the organic layer generating photons. In PCTWO90/13148 the organic light emissive material is a polymer. In US 4,539,507 the organic light emissive material is of the class known as small molecule materials, such as tris-(8-hydroxyquinolino)aluminium ("Alq3"). In a practical device, one of the electrodes is typically transparent, to allow photons to escape the device.

This technique for varying refractive index could be used for matrices of other materials, especially other organic materials. The organic materials could be polymers, oligomers or small organic materials or blends of two or more such materials. As illustrated above the refractive index of the resulting material could be fixed by appropriate choice of the material of the nanoparticles and their volume fraction in the matrix. The effective refractive index could be estimated by the Bruggeman effective medium approximation. The refractive index is thus adjustable depending on the refractive index of the organic matrix and the nanoparticle, the loading volume fraction, and the dispersion morphology.

A prior approach to adjusting the refractive index of organic materials has been to blend two or more organic materials together. However, the effectiveness of this approach has been limited because organic materials tend to have very similar refractive indices, and are often incompatible, leading to large-scale phase separation.

Example 2: Polymer distributed Bragg reflectors (Polymer mirrors)

A distributed Bragg reflector (DBR) consists of a stack of regularly alternating higher- and lower-refractive index dielectrics (light transmissive materials) fabricated to fulfil the Bragg condition for reflection at the design wavelength. This occurs when the optical path of the periodicity in the dielectric stack corresponds to half a wavelength, and the reflectivity is further optimised when the DBR stack obeys the following equation:

$$\frac{1}{2} \lambda = n_1 d_1 + n_2 d_2,$$

where n_1 , n_2 are the respective refractive indices; d_1 , d_2 are the corresponding component film thicknesses in the DBR; and λ is the design wavelength.

Figure 14 shows a polymer distributed Bragg reflector (polymer mirror) in which the alternate layers are formed of PPV and PPV modified by the dispersion of TiO_2 nanoparticles to tailor its refractive index. To form the polymer DBR a 0.4w/v% precursor PPV-MeOH solution (Cambridge Display Technology) containing 70ppm AOT as surface tension modifier and a 0.7w/v% (total solids) precursor PPV: TiO_2 -MeOH solution (in which the ratio of precursor PPV to TiO_2 is 1:1.8 by weight) were alternately spun on glass and thermally converted at 180°C under Ar for 1h to gave 600-700Å thick films of each material. The PPV material 10 forms the higher refractive index layers while the PPV: TiO_2 11 forms the lower refractive index layers.

The exact spin conditions which determine the thicknesses d_1 and d_2 were set after measurement of the reflection spectra upon the completion of each pair of high-and low-refractive index layers. Feedback is thus provided to allow a DBR to be engineered with the desired value of $n_1 d_1 + n_2 d_2$.

The spin coating could be performed on to a substrate of, for example, glass or plastic, which could in addition have a transparent conducting layer for charge injection.

Reflection spectra for three- and six-pair polymer layers with a first order reflection maximum at about 600nm (see figure 15) and 550nm (see figure 16) show the high level of control that can be achieved.

Since the DBR is formed of conjugated material it could be electrically-pumped to generate photons in addition to reflecting.

Example 3: Separate Confinement Heterostructures: Edge-emitting Organic Light Emitting Diodes

The ability to vary the refractive index of a material by dispersing nanoparticles in it provides an important enabling technology for the fabrication of photonic structures. An example is a photonic structure that makes use of refractive index contrasts (variations) to confine optical photon modes, for example to form waveguide structures and/or separate confinement heterostructures (see for example: S. M. Sze, "Semiconductor Devices: Physics and Technology," John Wiley & Sons, New York, 1985). The inability to conveniently vary refractive index has hitherto been a major obstacle to the full exploitation of organic materials in optoelectronic device technologies.

Figure 17a shows an edge-emitting organic light-emitting diode (EEOLED) that emits in the plane of the LED rather than vertically through it as in conventional surface emitting LEDs.

The EEOLED is fabricated by spin coating to give a multilayer structure as shown in figure 17a. There is a 3500Å thick low-refractive index layer of PPV:TiO₂ 12, followed by a 2500Å thick high-refractive index wave-guiding layer of PPV 13, and then a 1000Å thick emitter layer of MCP 14 which is fabricated with a number of thin charge-transport "hurdle" layers of PPV:SiO₂ 14a buried within the MCP layer. These PPV:SiO₂ thin layers provide a refractive index almost matching that of the adjacent MCP layers, but impose a weak electron and hole transport "hurdle" which can be exploited to enhance and direct recombination to occur largely in the MCP layer.

Hence, during operation of the device, excitons are produced in this layer. Light emitted from this layer is then wave-guided along the low-refractive index layer 12, which confines the photons, and emitted along the edge of the polymer film (see figure 17b). The excitons and photons are thus confined to separate regions of the device. In such a device the material properties of the region to which the excitons are confined can be tailored separately from those of the region to which the photons are confined: charge response and optical response properties of the conjugated material can be separated and individually modified or engineered to produce the desired electronic and photonic structure.

One advantage of this structure is that since the light is emitted from the edge of the device rather than the side there is no need for one of the electrodes of the device to be transparent, as in prior devices. Another advantage is that this structure forms the architecture for a wave-guiding electrically-pumped injection laser.

Example 4: Nanoparticles as a fluorescent dye carrier

By attaching functional materials to nanoparticles and dispersing those nanoparticles in a matrix the properties of the matrix can be adjusted. As an example of this technique, a device in which fluorescent dye is attached to nanoparticles and dispersed in a PPV matrix will be described.

A (sub)monolayer of the well-known red laser dye Rhodamine 101 is first adsorbed on to SiO₂ nanoparticles prepared as described above. To achieve this equal volumes (1ml) of a dilute 1w/v% solution of Rhodamine 101 (a Xanthene) in EtOH was mixed with 1.1w/v% SiO₂ in MeOH, and the homogeneous mixture dialysed against 200ml HPLC grade MeOH through a 6-8k MWCO cellulose dialysis tubing (Spectra/Por™) twice. Rhodamine 101 molecules not bound to the surface of the SiO₂ particles were dialysed out with the dialysate, whereas bound dye molecules were retained on the surface of the dispersed nanoparticles in the retentate. The retentate was then analysed by ultraviolet-visible spectrometry (see figure 18) to

illustrate the principle that the surfaces of nanoparticles could indeed be used as a dye carrier.

The nanoparticles prepared in this way could then be used as for the nanoparticles of example 1 to disperse the fluorescent dye into a layer of PPV. This layer could form the active layer of an organic light emitting device, in which the fluorescent dye could be stimulated by emission from the PPV to emit light at another frequency. This could be particularly significant in the manufacture of tri-colour displays.

Further examples of the surface modification of nanoparticle surfaces with fluorescent dye molecules are given in B. Oregan and M. Gratzel, "A low-cost, high efficiency solar-cell based on dye-sensitised colloidal TiO_2 films," *Nature* 353 (1991) pp. 737-740. The surface modification could be preceded by surface modification by a suitable chemical, such as a silylation reagent, to cap the nanoparticles' surfaces with a layer carrying appropriate chemical functional groups (such as $-\text{COOH}$, $-\text{NH}_2$, $-\text{SO}_3\text{H}$, $-\text{PO}_3\text{H}$ or $-\text{NR}_3^+$) on which covalent or ionic or hydrogen bonds could be formed with an appropriate functional group on the dye molecule. This could help the dye to anchor to the surface of the nanoparticles. This may require modification of the dye molecules themselves to produce the desired functional group.

A wide range of fluorescent dye molecules are available, to provide sensitised emission over a wide range of frequencies, especially when used as a "guest" in a "guest-host" active layer structure where the host is a light emissive material which can stimulate the guest. In these structures, the matrix "host" layer, which could be of an organic material, performs the role of charge transport and acts as first-stage recombination centres, but subsequently the energy is to be transferred to the "guest" which then emits the desired colour. It is preferred that the energy transfer from the initial excitation created in the matrix to the final excitation to be created in the dye should be efficient, which in general demands that the emission spectrum of the initial ("host") excited state should overlap significantly with the absorption spectrum to the final ("guest") excited state. It is also preferred that the guest should exhibit a high photoluminescent efficiency: and that the density of the "guest" sites in the

"host" matrix should be sufficiently high (preferably 10^{-17} to 10^{-19} /cm³) if the effective radius of energy transfer (the Forster radius) is small, e.g. about 3-10nm.

Example of suitable dyes may include the class of molecules widely known as the laser dyes (such as coumarins, xanthenes and oxazines).

In the past a major stumbling block to the incorporation of dye molecules in matrices such as organic materials has been rapid diffusion and segregation (recrystallisation) of the dye molecules in the host layer or at the interfaces during storage, and especially during operation, so that emission efficiency falls rapidly. This problem can be addressed by the process described above of immobilising the dye by binding it to the surface of a nanoparticle which is relatively immobile in the matrix. Even with a diameter of 1-10nm, the diffusion coefficient of a nanoparticle through a typical organic host could be expected to be several orders of magnitude lower than that of the dye molecules themselves.

Example 5: Nanoparticles as a Polymer Morphology Modifier

This example demonstrates that nanoparticles that are intimately dispersed in a polymer matrix can modify the morphology (e.g. the chain conformation) of the polymer.

Polymer films of PPV:TiO₂ (having the same composition as the film C described above) and PPV (as control) were spun and subjected to atomic force microscopy examination. Both films were thermally converted at 180°C. Figures 19a and 19b show the results of the examination. The surface of the control film exhibits a number of nanoscale protruding domains (about 10-20nm high and 20-40nm wide) that are believed to be related to micro-crystallisation of the PPV chains in the film thickness direction (M. A. Masse, D. C. Martin, E. L. Thomas and F. E. Karasc, "Crystal morphology in pristine and doped films of poly(p-phenylene vinylene)," Journal of Materials Science 25 (1990), pp. 311-320). In contrast, films fabricated with the nanoparticles showed a relatively smooth featureless surface over the same

length scale. This is believed to be due to the suppression of micro-crystallisation by the nanoparticles, probably by steric constraints and the introduction of kinks in the chain conformations in the interfacial layer next to the nanoparticle surface.

Further evidence for this belief is given by ultraviolet-visible spectrometry of a set of films with a range of PPV:TiO₂ volume ratios, which showed a progressive blue shift of the π - π^* transition with increasing nanoparticle content above 15% (see figure 20).

Thus, it appears that the presence of nanoparticles can induce perturbation of local polymer morphology, changing the effective conjugation length and/or crystallinity of the polymer matrix and thus its optical refractive index and other related properties.

This feature of organic/nanoparticle composites could be used to inhibit the recrystallisation of the organic during storage or in operation. Recrystallisation is generally undesirable as, taking optical devices as an example, it can cause changes in electrical behaviour, colour stability and luminous efficiency, with a deleterious impact on device performance. Altering morphology could also alter many "intrinsic" properties such as transport, binding energies (as for excitons) and efficiency.

General Issues

Nanoparticles for the above applications and other applications could be formed of materials other than SiO₂ or TiO₂. Depending on the application some properties of the particles that may have to be borne in mind are:

1. Transparency. To achieve this the particles would have to have a sufficiently large band-gap. For instance, where the nanoparticles are dispersed in a matrix of an emissive polymer and are required to be transparent to emissions from that polymer the particles should have a larger band gap than the polymer material. One possibility is to use transparent, non-metallic, inorganic particles. It is preferred that the optical bandgap of the nanoparticle material is equal to or larger than that of the matrix material, so that the absorption edge is not squeezed into the optical transparency window.

2. Insolubility (e.g. in water). This could be assessed in the first approximation by behaviour towards water as detailed in the CRC Handbook.
3. Resistance to acid corrosion. This is especially relevant where the nanoparticles are to be used in an acid environment, e.g. in PPV blends. This property could be assessed in the first approximation by bulk behaviour towards inorganic acids as detailed in the CRC Handbook.
4. Non-mobile ion forming. A cation-exchange process could be used to remove unwanted ions such as Na^+ from the preparation if they would otherwise interfere with performance.
5. Size. If uniform charge transport and high optical quality of the resulting body is a requirement then a preferred size range is about 5-10nm. Particles greatly smaller than this may be undesirable because of lowered stability due to their higher surface-to-volume ratio. For transparency of bodies containing the nanoparticles it is preferred that the nanoparticles' diameters are generally less than half of the relevant light wavelength. If the body is small, such as a typical thin film of an optical device, it is preferred that the nanoparticles' diameters are less than half the thickness of the film, most preferably less than one fifth or one tenth of the thickness of the film. For a uniform (statistical) distribution the size of the nanoparticles should be 1/5 (1/10) of the relevant size (wavelength or film thickness).
6. Dispersion. To improve dispersion of the nanoparticles, and reduce agglomeration, a surfactant coating could be provided on the surface of the nanoparticles. One example of a potentially suitable surfactant is sodium decussate (AOT). This is an ionic surfactant with short hydrophobic tails in the form of 2-ethylhexyloxy ester side-chains that may be expected to be compatible with and hence dispersible in MEH-PPV and a variety of polar casting solvents like MeOH and CHCl_3 .
7. Suitable refractive index.
8. Processable into homogenous blends with the matrix material, for example a desired emissive polymer.

Nanoparticles may be "blended" with sublimed molecule materials by forming a dispersed layer of particles and then subliming molecules on top of that layer. In this

way the molecules (oligomers etc.) can fill in the gaps set by the particles. The formation of the first layer could be through spin coating or self assembly. This technique may also allow nanoparticles to be blended with materials that can be deposited by evaporation at high temperature as inorganic materials are deposited in a CVD (chemical vapour deposition) or MBE process or any variant or combination.

The present invention may include any feature or combination of features disclosed herein either implicitly or explicitly or any generalisation thereof, irrespective of whether it relates to the presently claimed invention. In view of the foregoing description it will be evident to a person skilled in the art that various modifications may be made within the scope of the invention.

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CLAIMS

1. A method for preparing nanoparticles for use, from a mixture of nanoparticles with another material, the method comprising washing the mixture with a solvent in which the nanoparticles are soluble to remove the said other material and form a solution of nanoparticles in the solvent.
2. A method as claimed in claim 1, comprising separating at least a first fraction of the nanoparticles from a mixture of the solvent and the said other material.
3. A method as claimed in claim 2, wherein the separation is performed by filtration.
4. A method as claimed in claim 2, wherein the separation is performed by dialysis.
5. A method as claimed in claim 2, wherein the separation is performed by centrifugation.
6. A method as claimed in any of claims 2 to 5, wherein in the separation step the said first fraction of the nanoparticles are separated from another fraction of the nanoparticles.
7. A method as claimed in claim 6, wherein the nanoparticles of the said other fraction are relatively small in comparison to the nanoparticles of the first fraction.
8. A method as claimed in any preceding claim, wherein the said other material is a by-product of the formation of the nanoparticles.
9. A method as claimed in any preceding claim, wherein the said other material comprises a surfactant.
10. A method as claimed in any preceding claim, wherein the solvent is an organic solvent.

11. A method as claimed in any preceding claim, wherein the solvent is an alcohol.
12. A method as claimed in any preceding claim, wherein the said other material is soluble in the solvent.
13. A method as claimed in any preceding claim, wherein the solvent is one in which the said other material is preferentially soluble to the nanoparticles.
14. A method as claimed in any preceding claim, wherein the solvent is a polar solvent.
15. A method as claimed in any preceding claim, wherein the nanoparticles are generally smaller than 50nm in diameter.
16. A method as claimed in any preceding claim, wherein the solvent is such as to hold the dissolved nanoparticles in a disaggregated state.
17. A method as claimed in any preceding claim, comprising adding a surface modifying agent to the solution of nanoparticles.
18. A method as claimed in any preceding claim, wherein the surface modifying agent is a dye.
19. A method as claimed in any preceding claim, wherein the nanoparticles are light transmissive.
20. A method as claimed in any preceding claim, wherein the nanoparticles are non-conductive.
21. A method as claimed in any preceding claim, comprising adding the solution of nanoparticles to a polymer precursor.

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106040-12924260

22. A method as claimed in claim 21, comprising converting the polymer precursor to form a polymer body containing a substantially uniform dispersion of nanoparticles.
23. A method as claimed in claim 21 or 22, comprising treating the polymer to render it insoluble in the solvent.
24. A solution of nanoparticles formed by a method according to any of claims 1 to 20.
25. A polymer precursor material containing nanoparticles, formed by a method according to claim 21.
26. A polymer material comprising a substantially uniform dispersion of nanoparticles formed by a method according to any of claims 1 to 20.
27. A polymer material containing a substantially uniform dispersion of nanoparticles, formed by a method according to claim 22 or 23.
28. An organic material containing a substantially uniform dispersion of nanoparticles.
29. An organic material as claimed in claim 28, wherein the organic material is a semiconductive material.
30. An organic material as claimed in any of claims 26 to 29, wherein the presence of the nanoparticles influences at least one material property of the organic material.
31. An organic material as claimed in claim 30, wherein the said property is an optical and/or an electronic property.

32. An organic material as claimed in claim 31, wherein the said property is refractive index.

33. An organic material as claimed in any of claims 26 to 32, wherein the nanoparticles have a surface coating.

34. An organic material as claimed in claim 33, wherein the surface coating is of a material that influences at least one optical and/or electrical property of the organic material and/or influences the interaction of the nanoparticles with the organic material.

35. A method for tailoring at least one property of an organic material, the method comprising forming a substantially uniform dispersion of nanoparticles in the organic material.

36. A method for forming a solution of nanoparticles substantially as herein described with reference to the accompanying drawings.

37. Organic material containing a substantially uniform dispersion of nanoparticles substantially as herein described.

Energy dispersive x-ray spectrometry
Excitation, 20keV
Sample, thin TiO_2 particle film on carbon tape

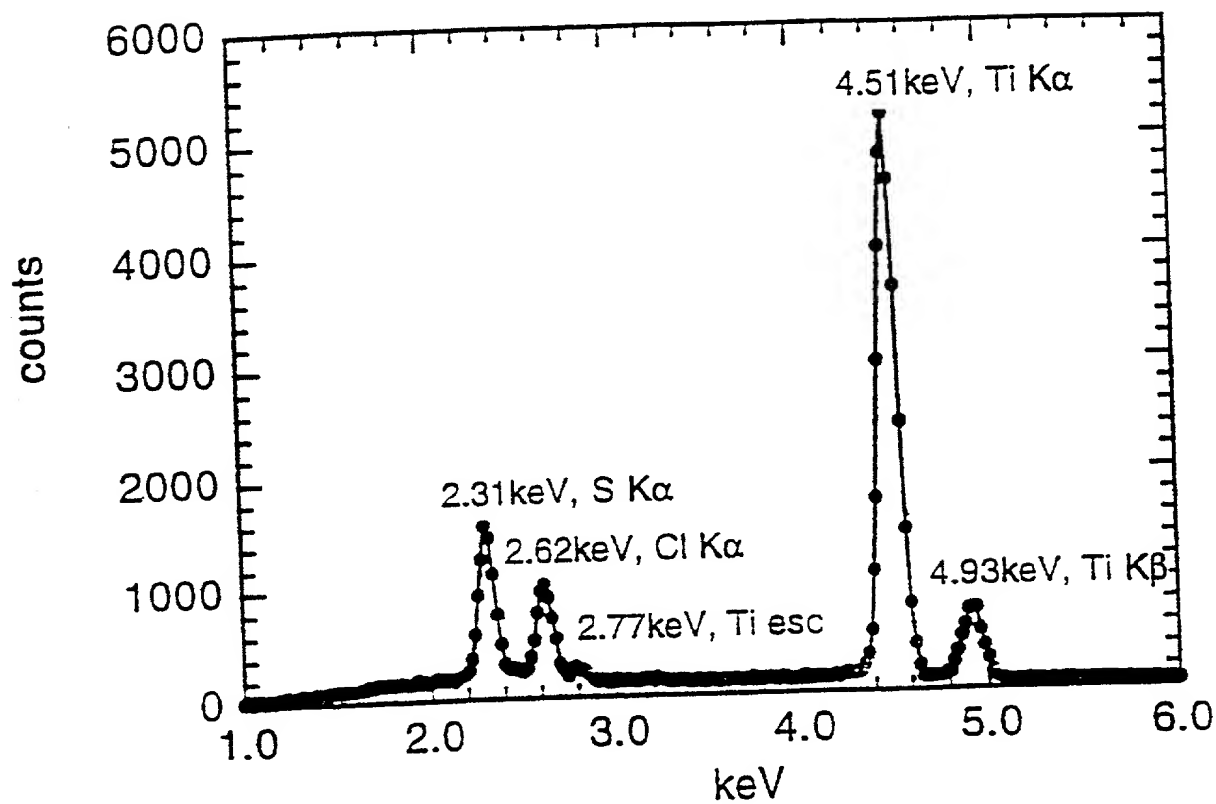


FIG.1

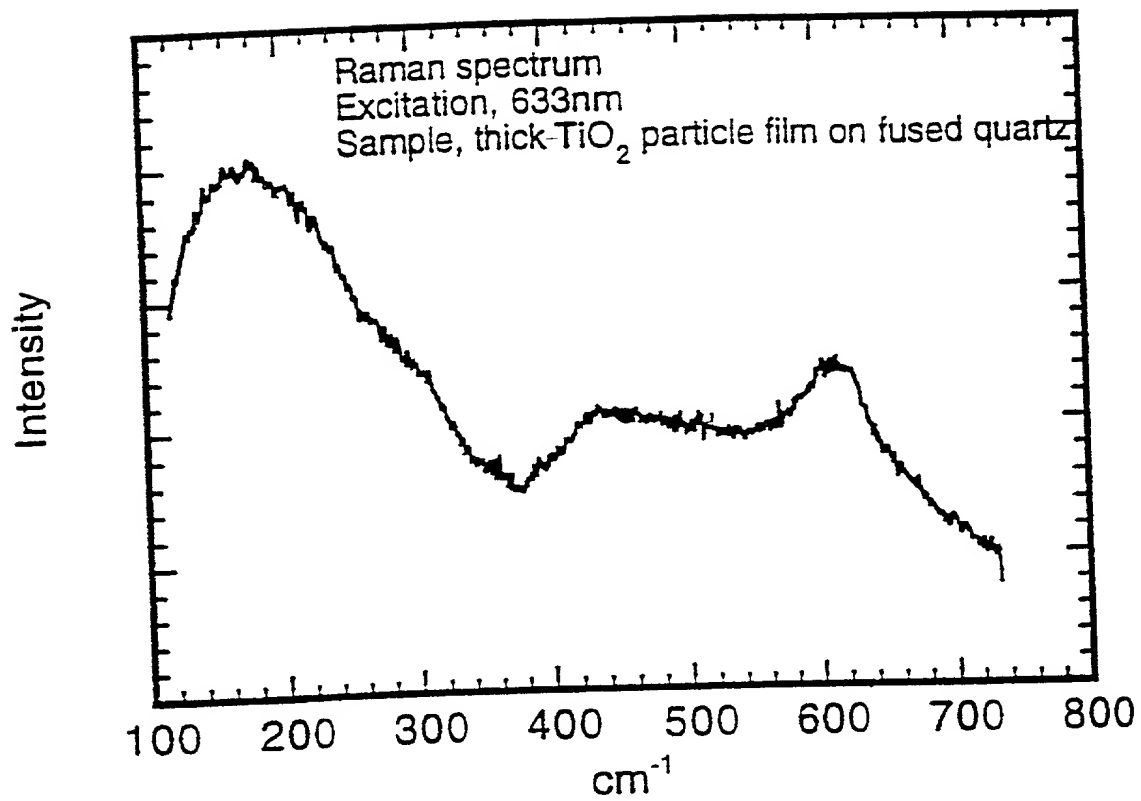


FIG.2

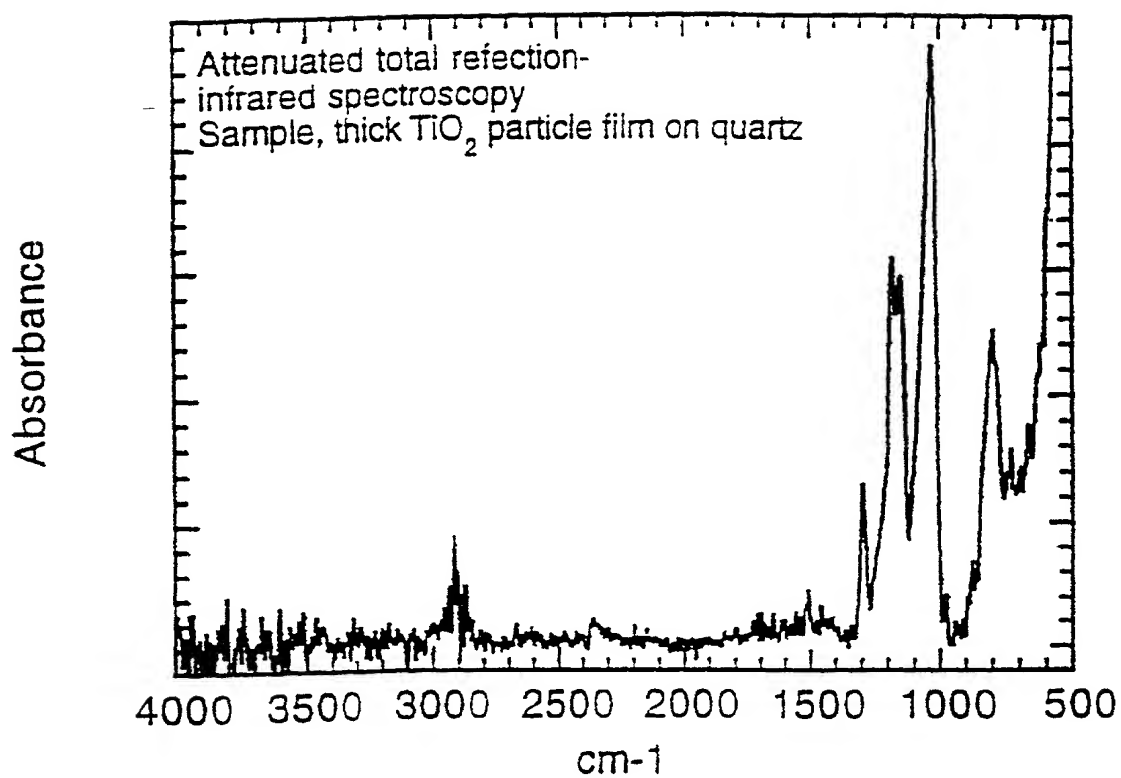


FIG.3

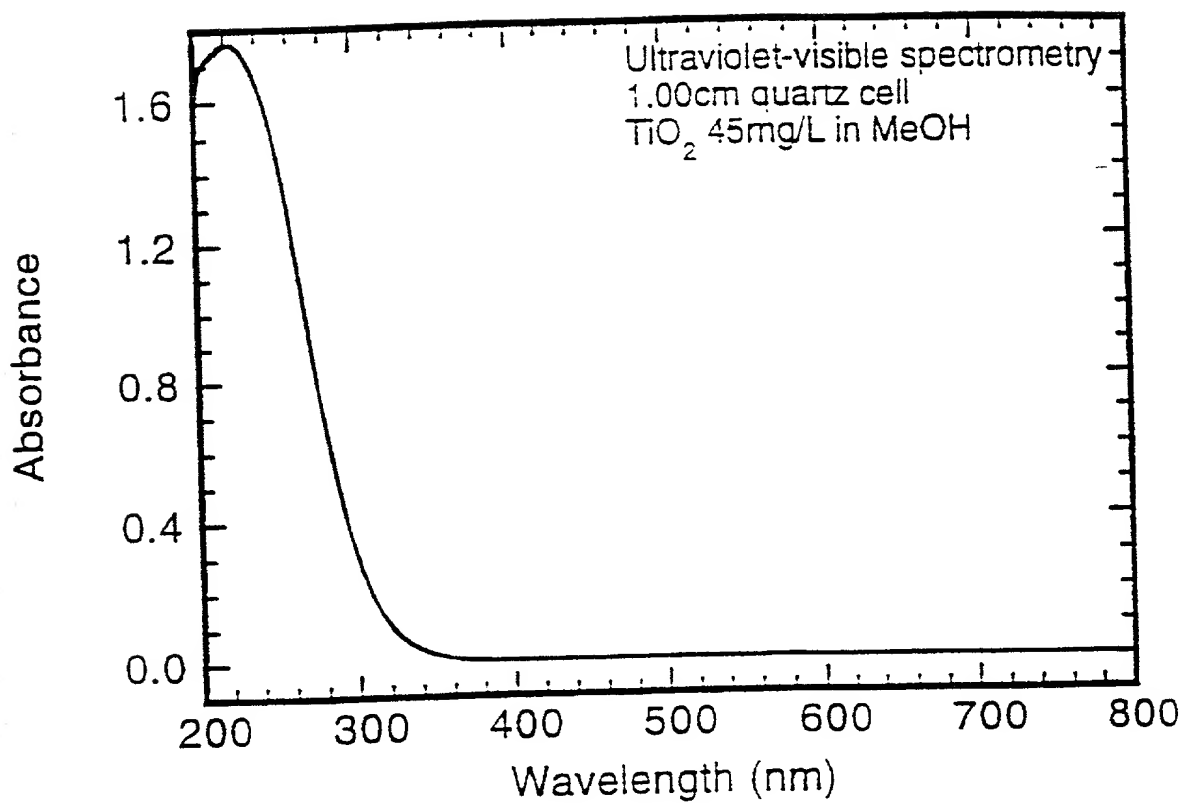


FIG.4



FIG.5

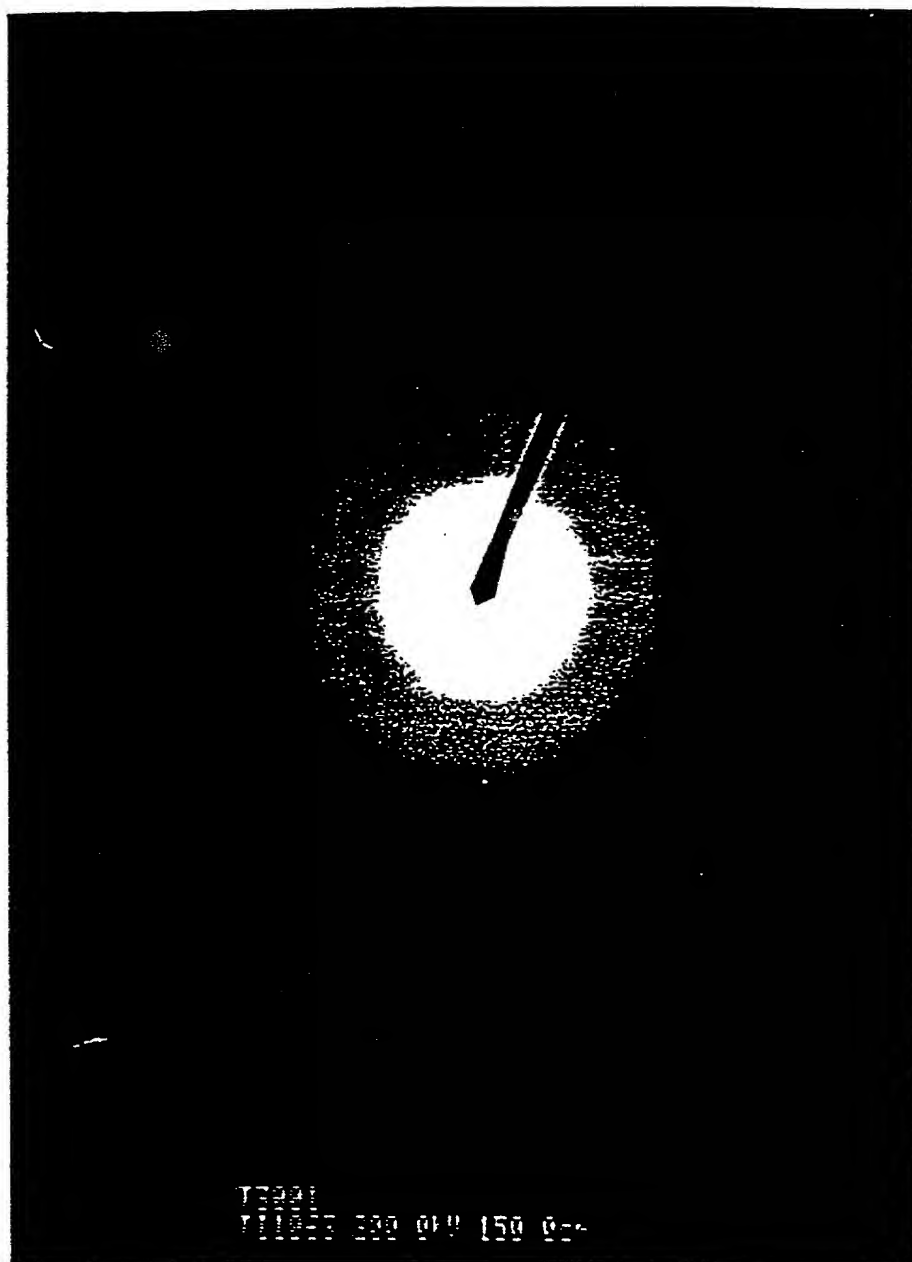


FIG.6

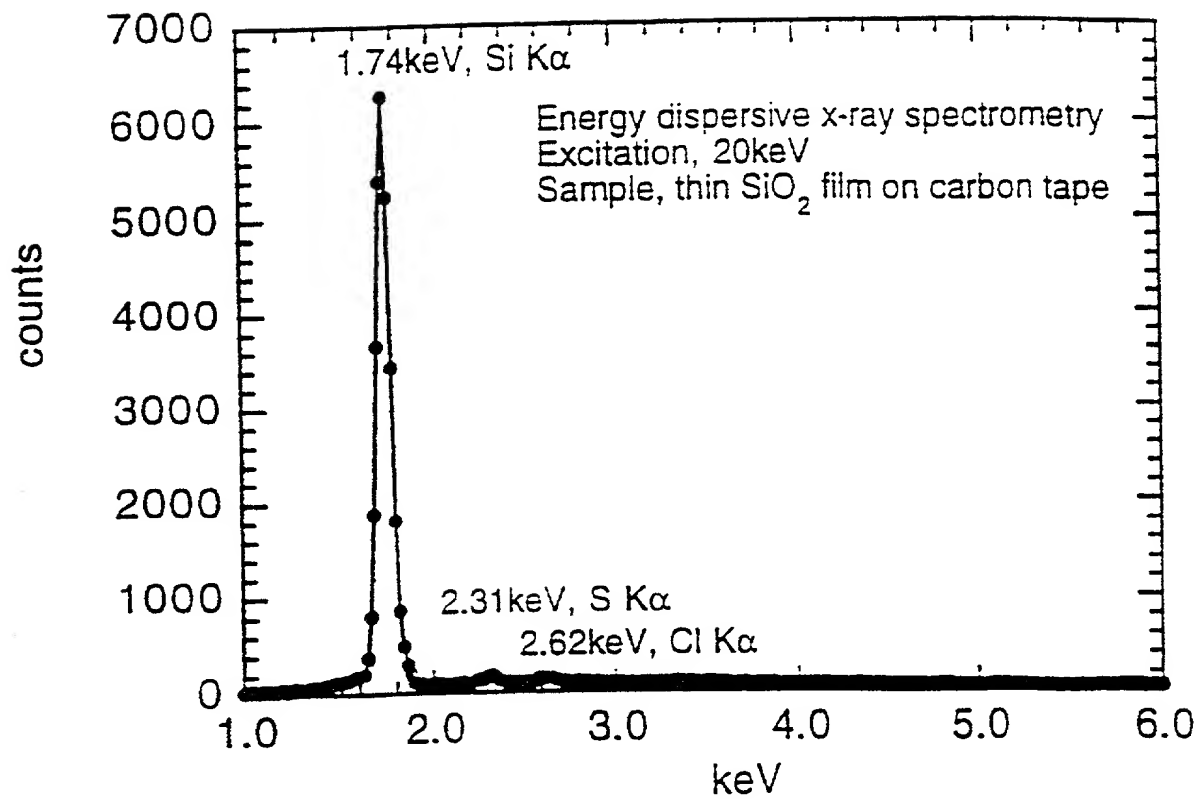


FIG.7

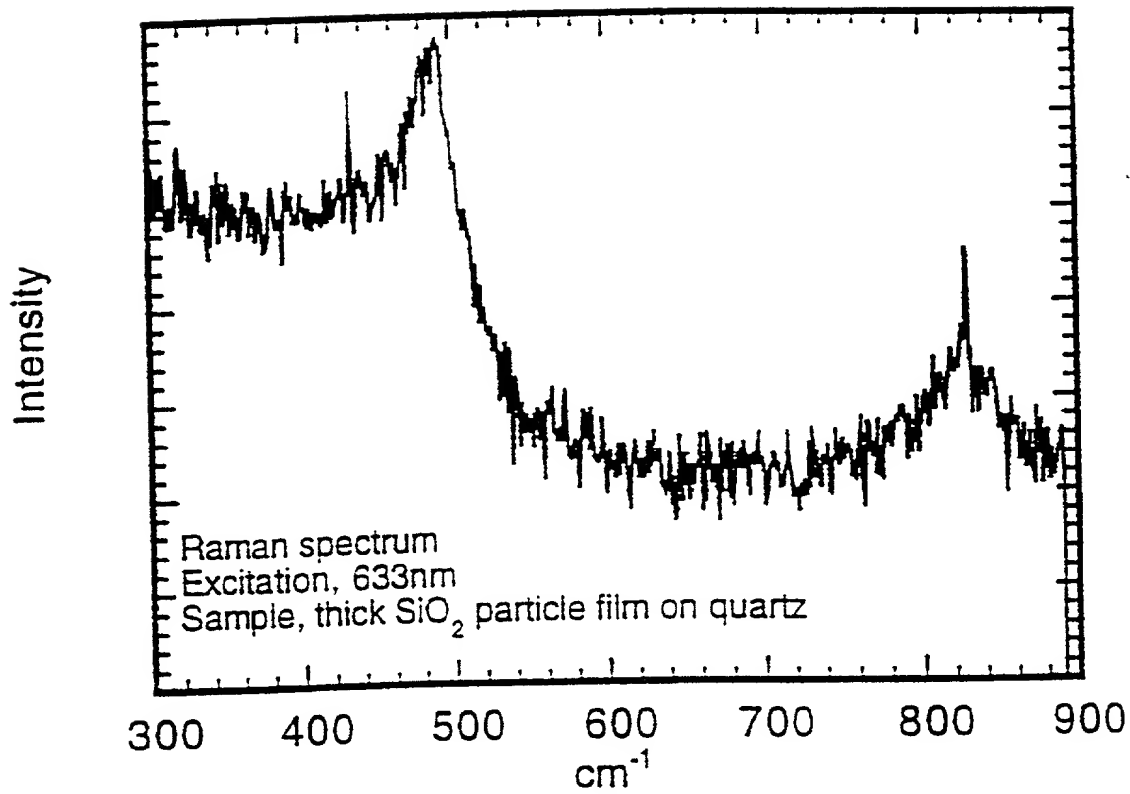


FIG.8

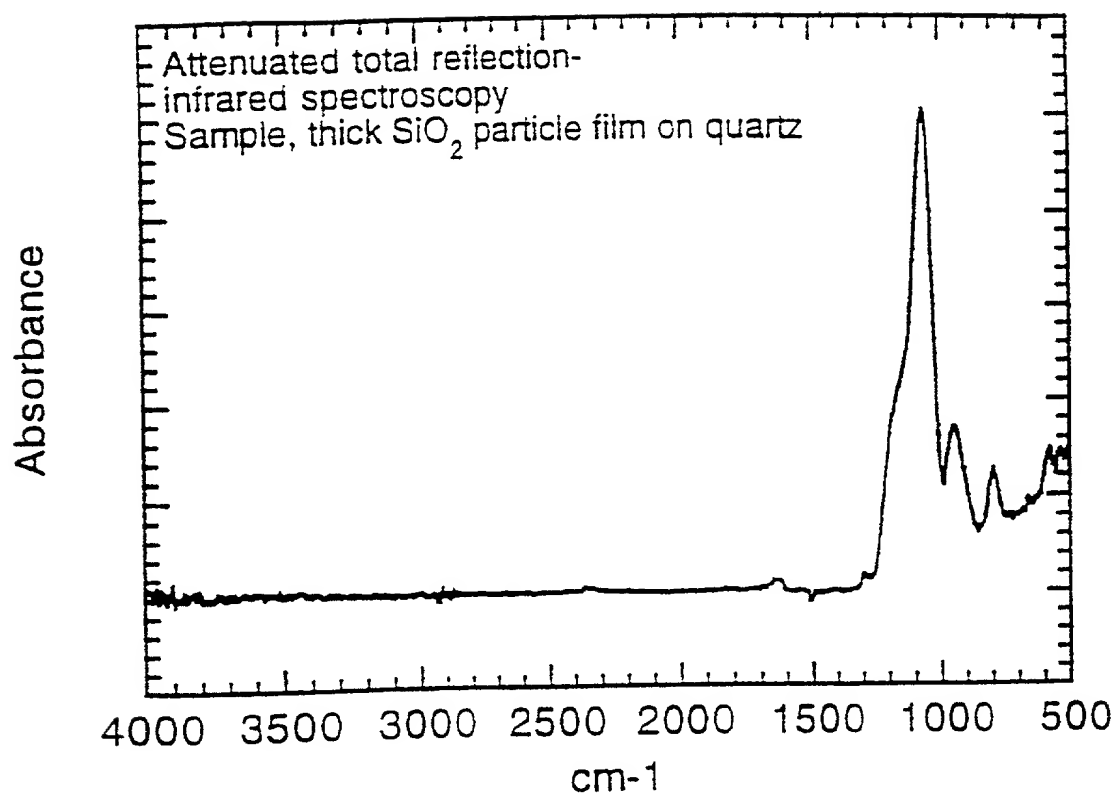


FIG.9



FIG.11

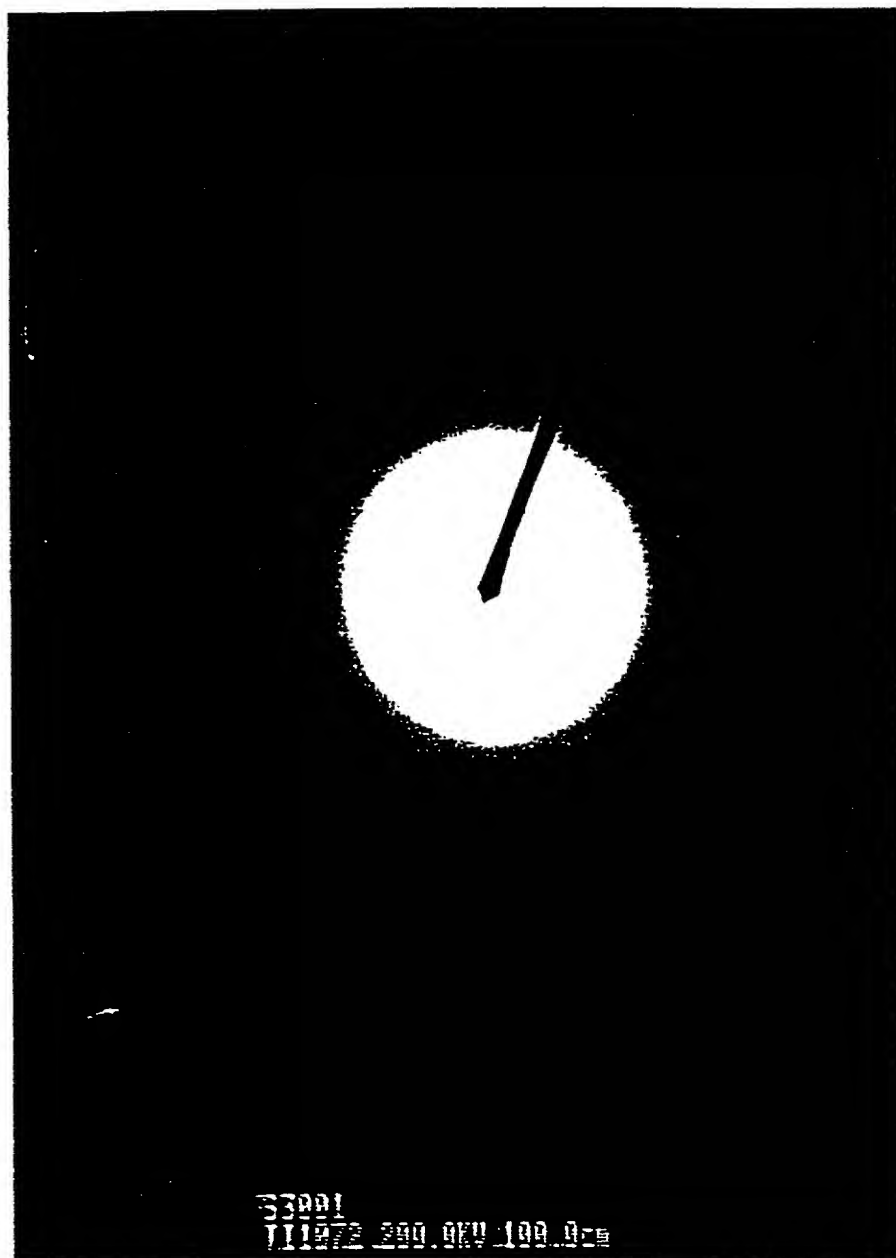


FIG.12

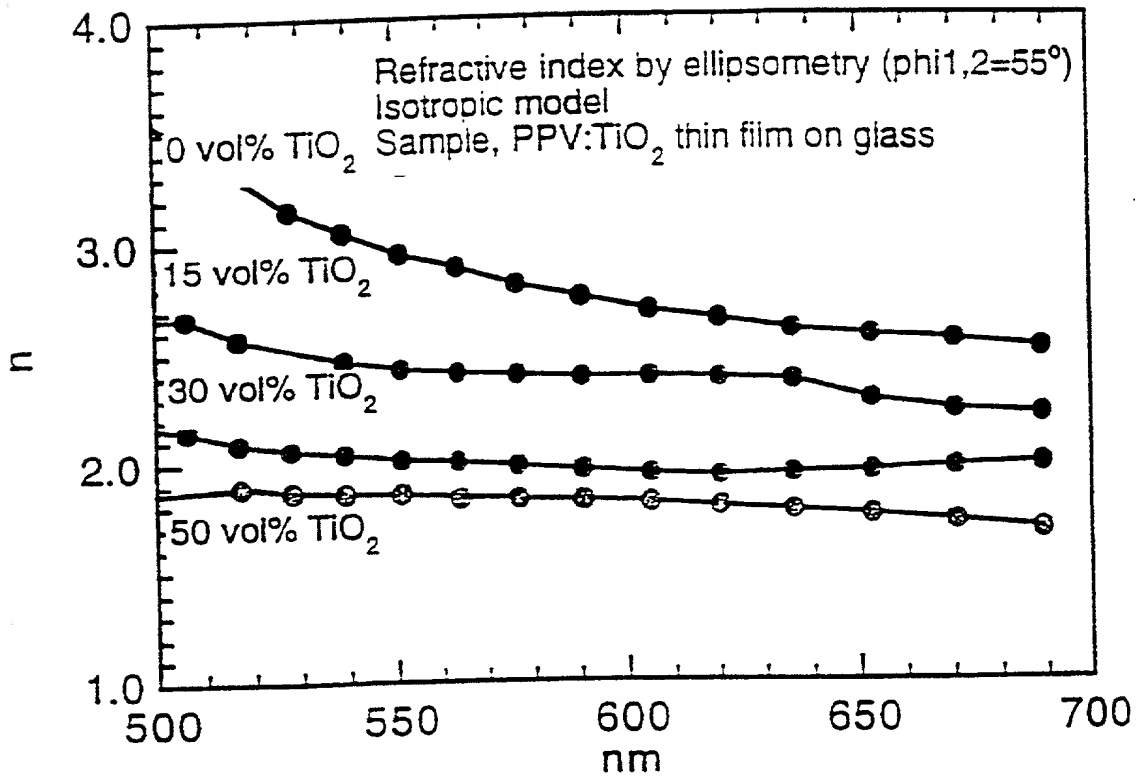
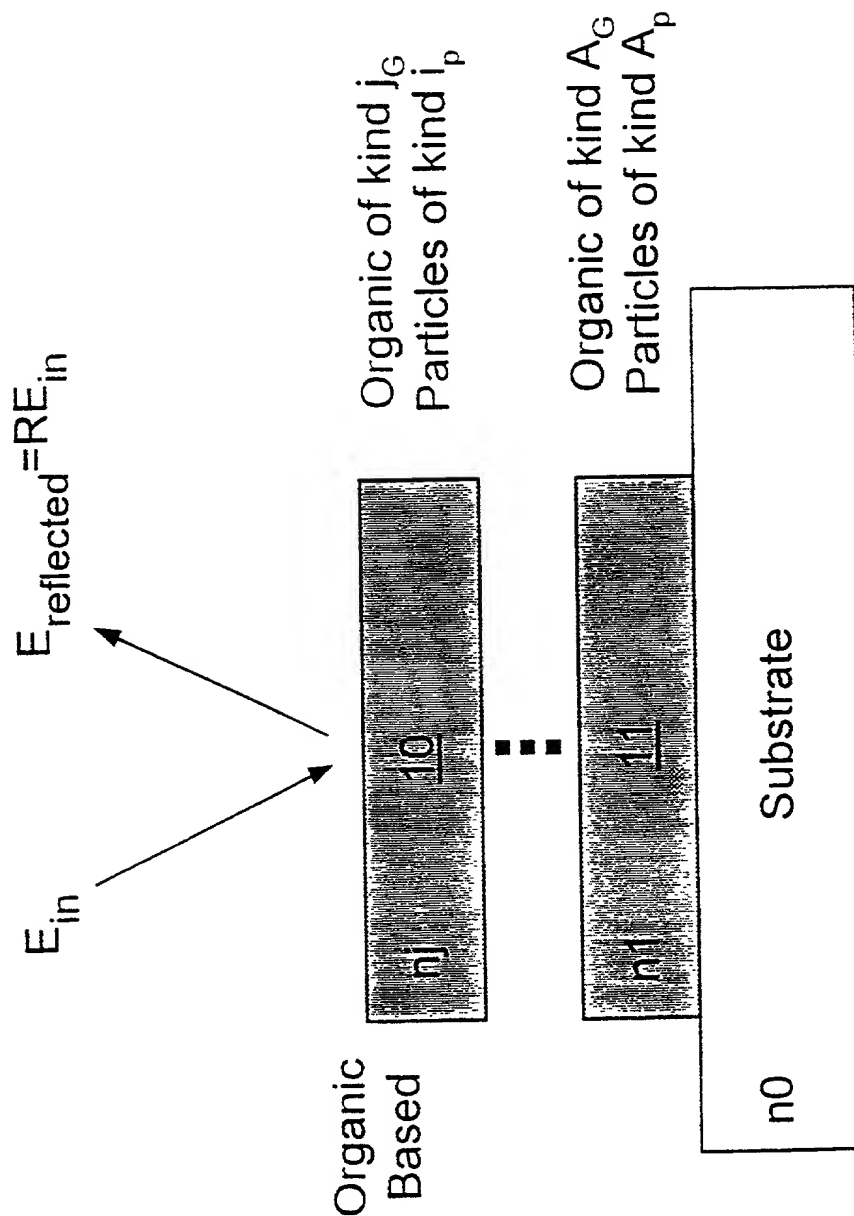


FIG.13

Distributed Brag Reflector



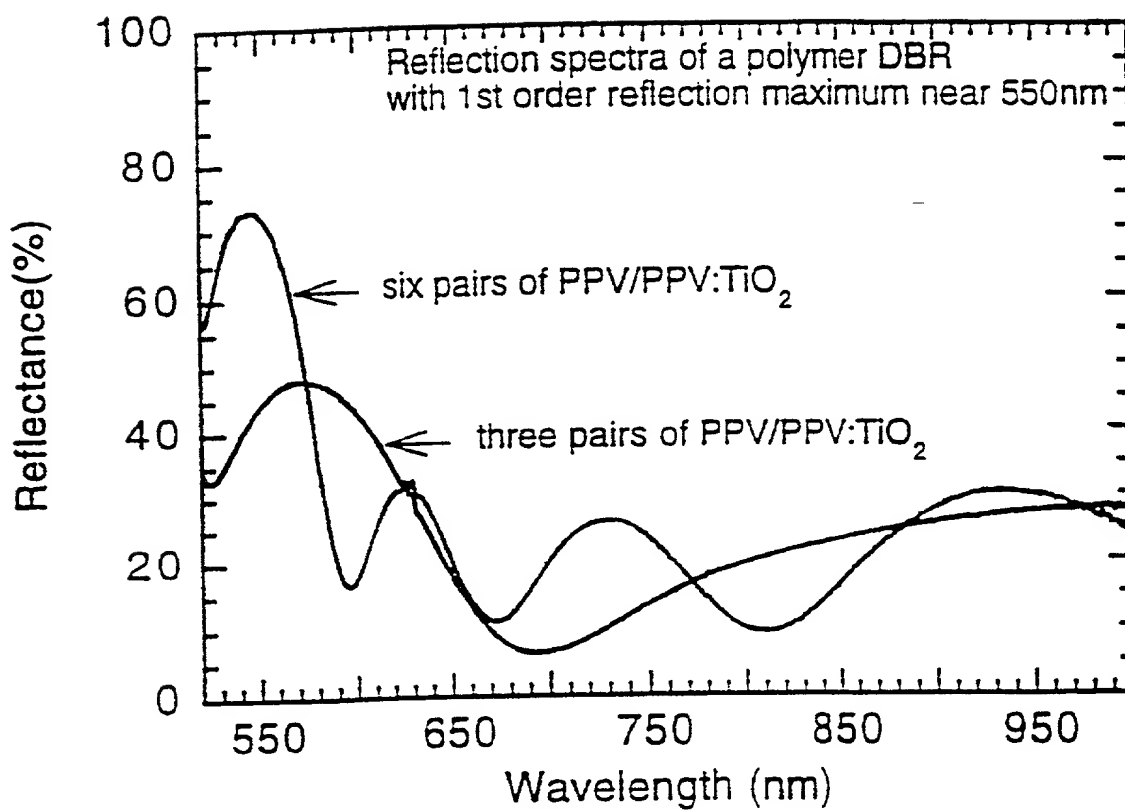


FIG.15

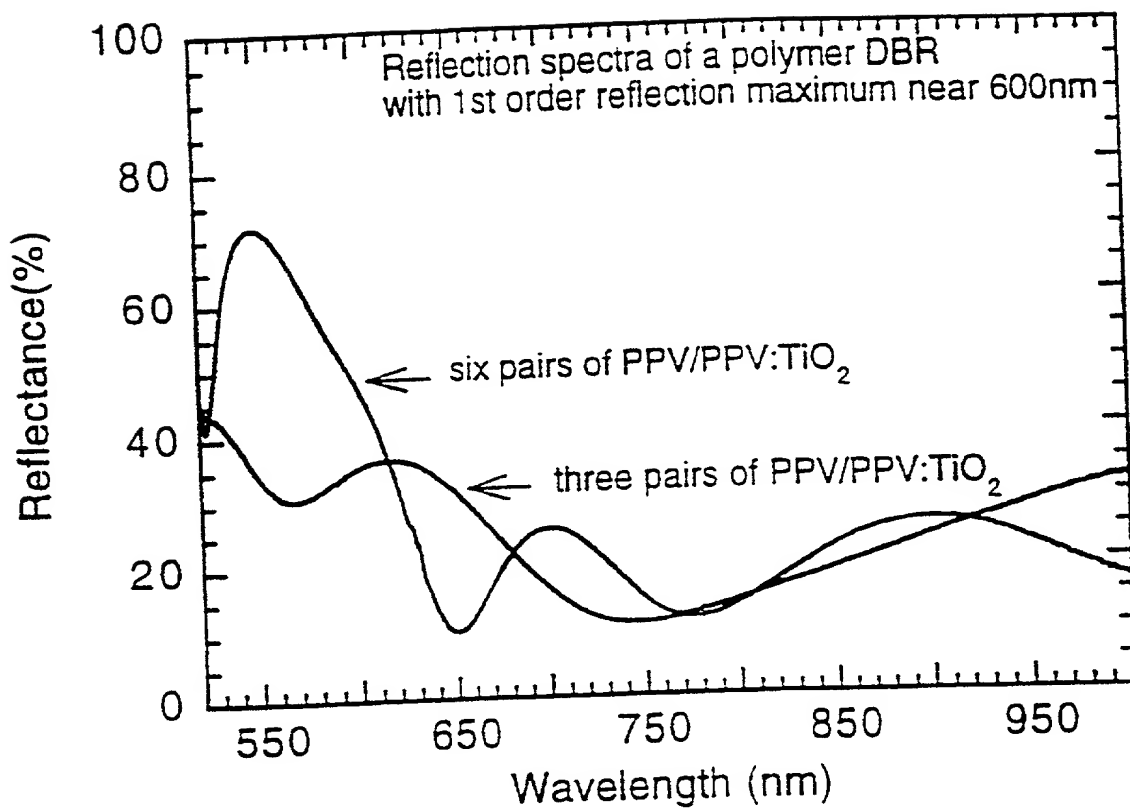


FIG.16

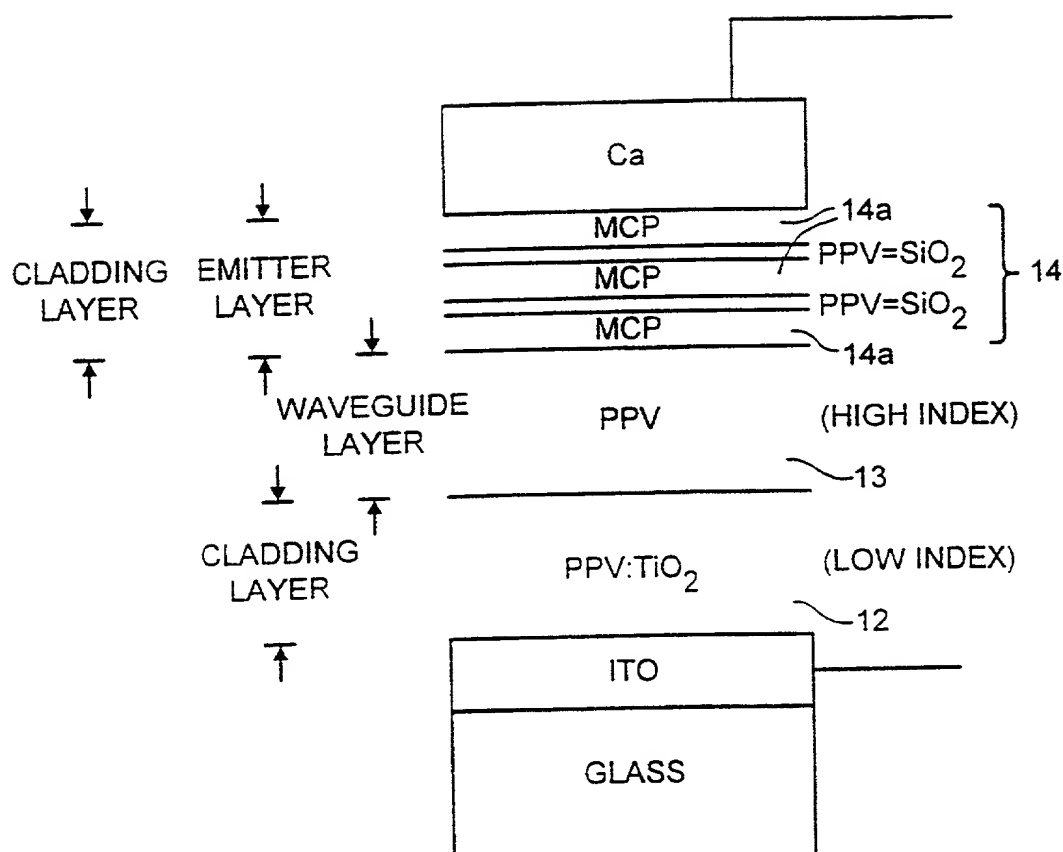


FIG. 17a

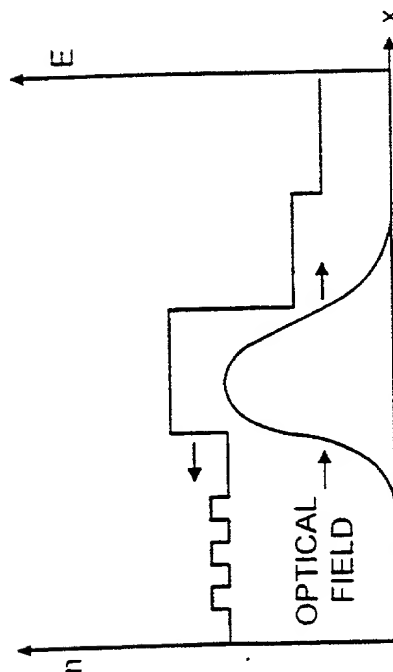
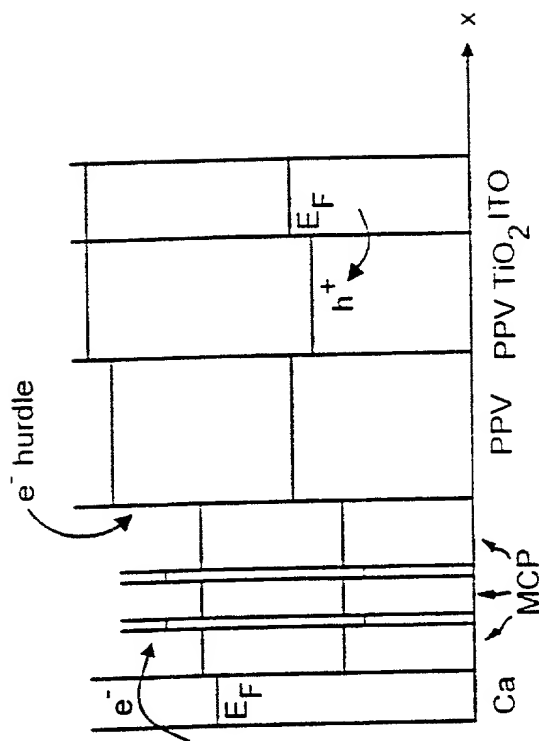


FIG. 17b

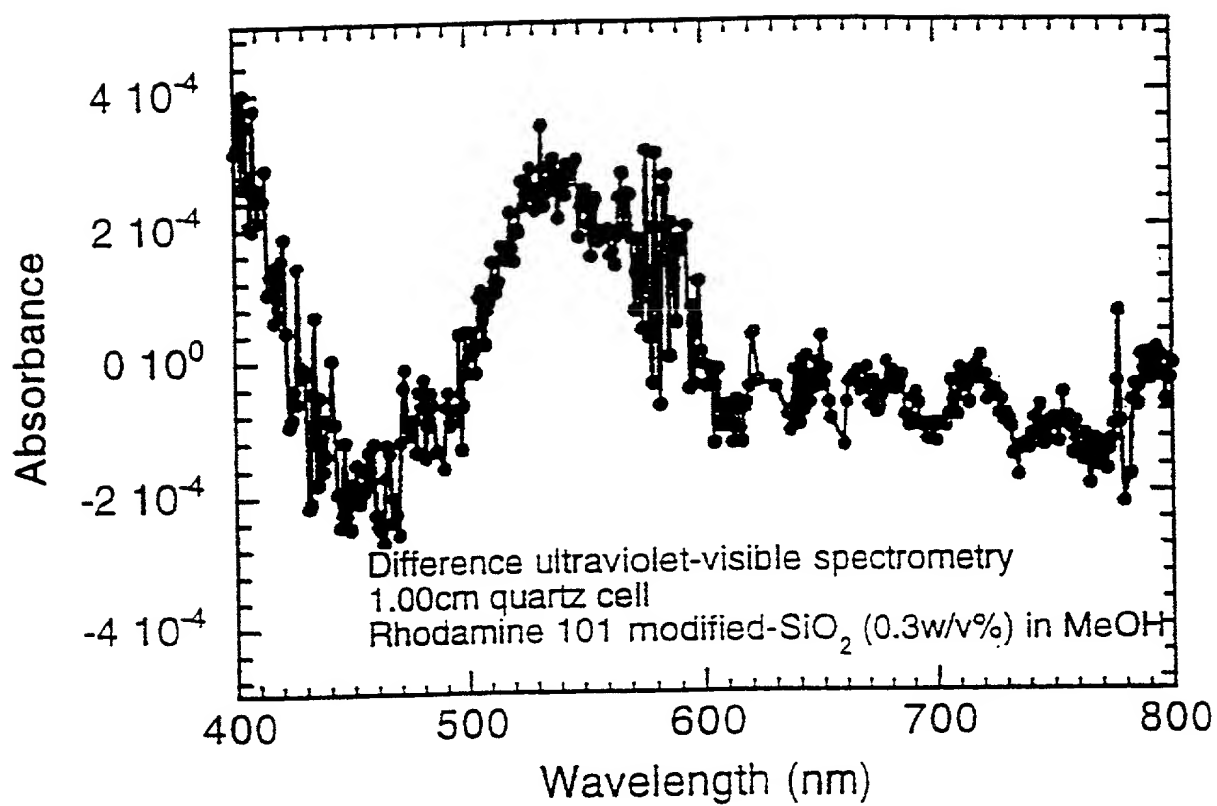
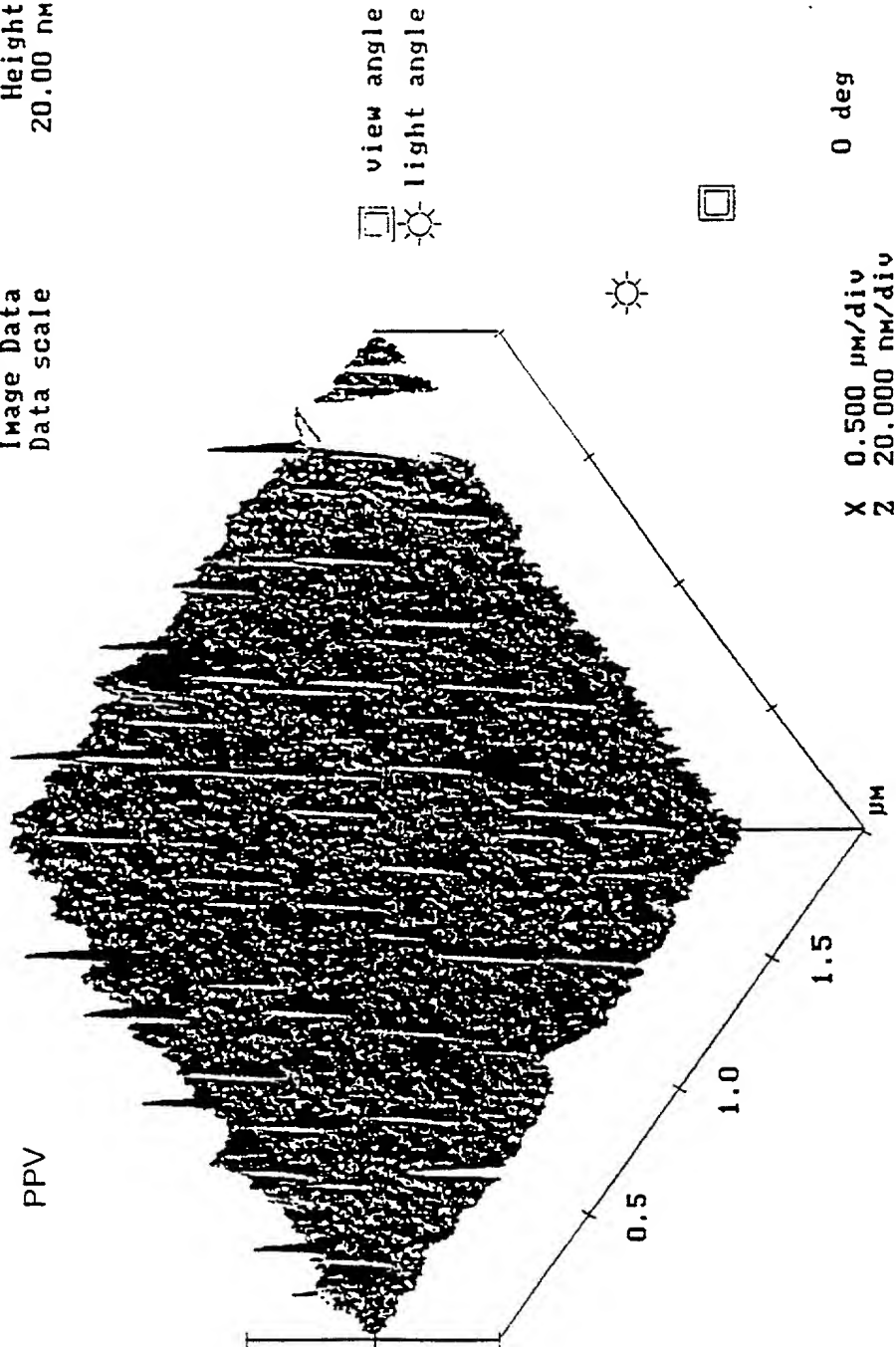


FIG.18

Digital Instruments Nanoscope
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Scan rate 1.001 Hz
Number of samples 512
Image Data Height
Data scale 20.00 nm

FIG. 19a

PPV

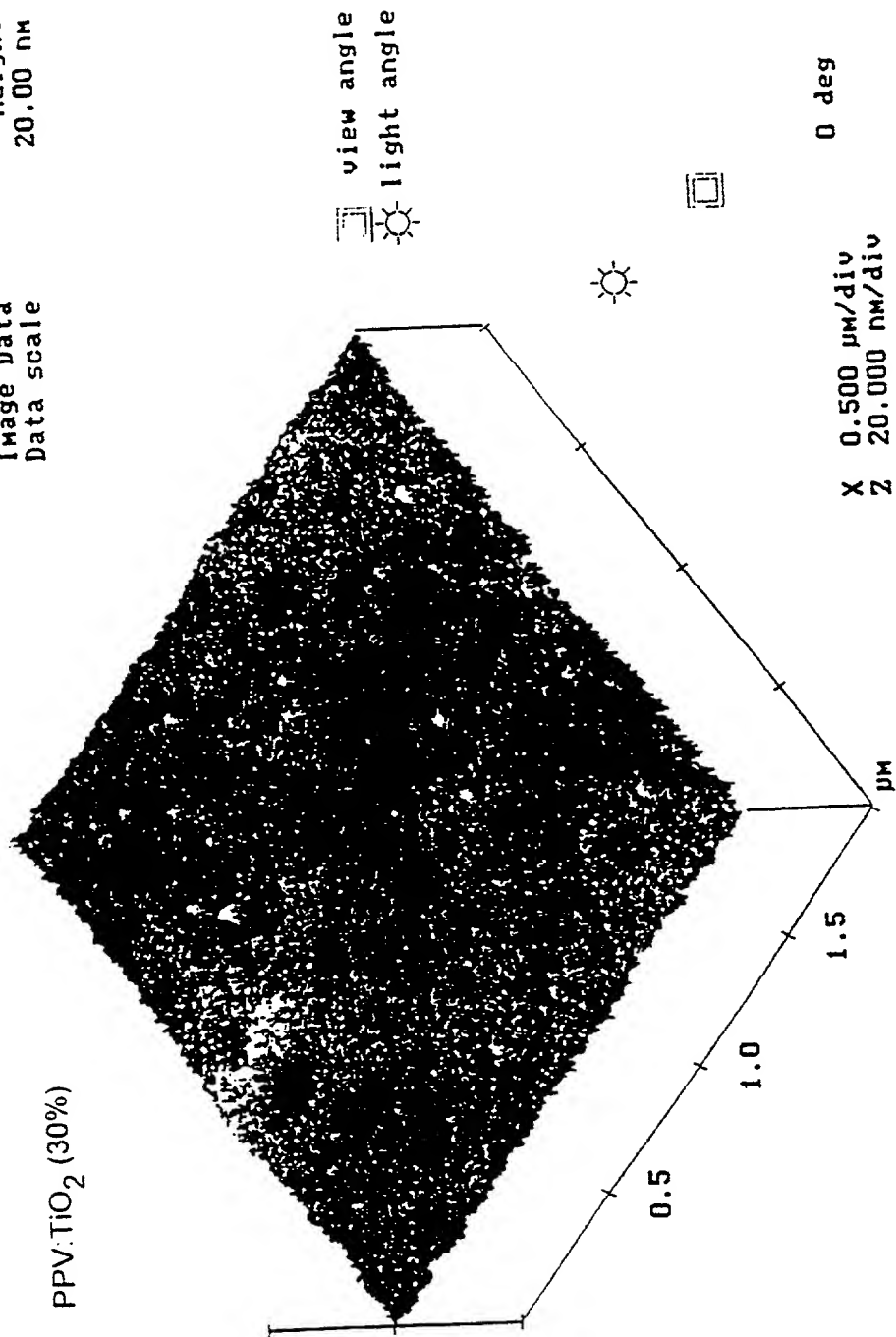


0-tppv.010

Digital Instruments Nanoscope
 Scan size 2.000 μm
 Scan rate 1.001 Hz
 Number of samples 512
 Image Data Height
 Data scale 20.00 nm

FIG. 19b

PPV:TiO₂ (30%)



30-tppv.010

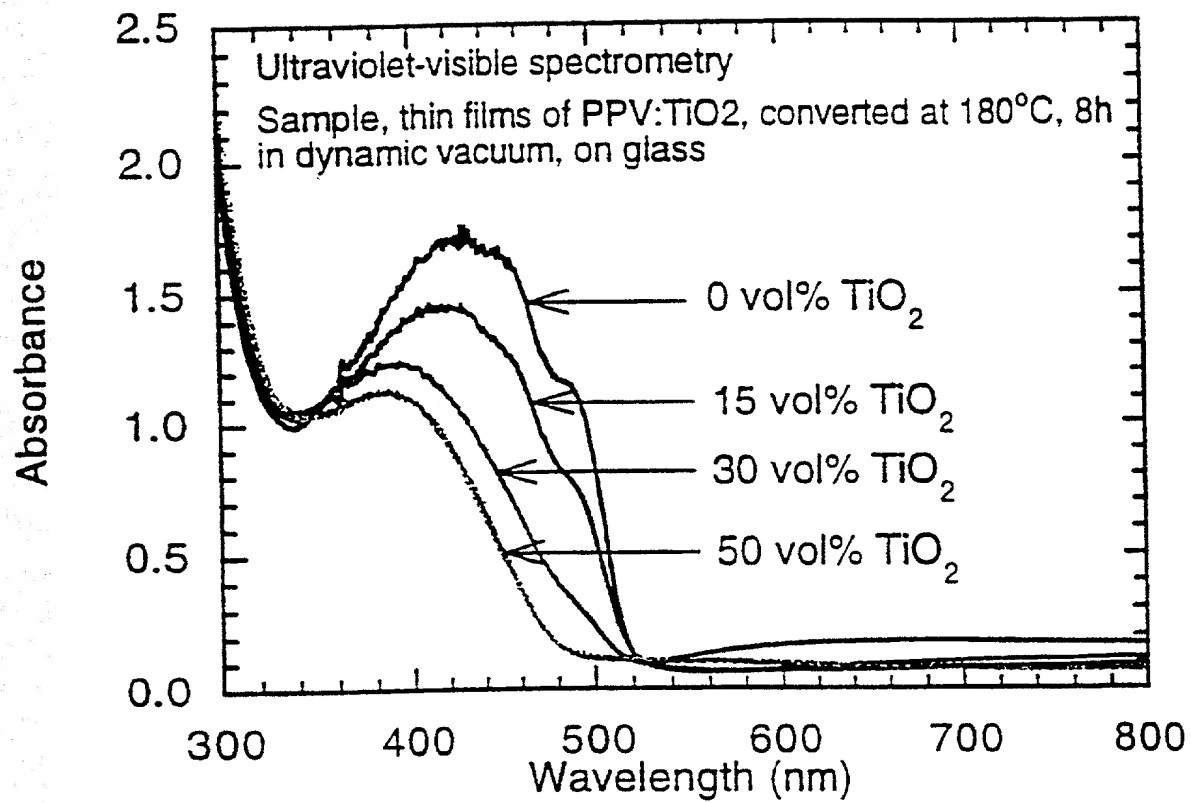


FIG.20



Attorney Docket No. C1043/7030

DECLARATION FOR PATENT APPLICATION

As a below named inventor, I hereby declare that:

My residence, post office address and citizenship are as stated below next to my name.

I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled

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the specification of which is attached hereto unless the following is checked:

January 12, 2001

☒ was filed on 09/743,621, as United States Application No. 09/743,621 or PCT

(Include Series Code)

International Application No. _____, bearing attorney docket No. C1043/7030, and was amended on _____ (if applicable).

I hereby state that I have reviewed and understand the contents of the above identified specification, including the claims, as amended by any amendment referred to above.

I acknowledge the duty to disclose information which is material to the examination of this application in accordance with Title 37, Code of Federal Regulations, §1.56.

I hereby claim foreign priority benefits under Title 35, United States Code, §119(a)-(d) or §365(b) of any foreign application(s) for patent or inventor's certificate. or section 365(a) of any PCT International application designating at least one country other than the United States listed below and have also identified below any foreign application for patent or inventor's certificate or PCT International application having a filing date before that of the application on which priority is claimed:

Prior Foreign PCT International Application(s) and any priority claims under 35 U.S.C. §§119 and 365 (a), (b):

(Number)	(Country-if PCT, so indicate)	(DD/MM/YY Filed)	Priority Claimed
9815270.5	GB	14 July 1998	<input checked="" type="checkbox"/> YES <input type="checkbox"/> NO
(Number)	(Country-if PCT, so indicate)	(DD/MM/YY Filed)	<input type="checkbox"/> YES <input type="checkbox"/> NO
(Number)	(Country-if PCT, so indicate)	(DD/MM/YY Filed)	<input type="checkbox"/> YES <input type="checkbox"/> NO
(Number)	(Country-if PCT, so indicate)	(DD/MM/YY Filed)	<input type="checkbox"/> YES <input type="checkbox"/> NO

I hereby claim the benefit under Title 35, United States Code, §119(e) of any United States provisional application(s) listed below:

(Application Number)	(filing date)
(Application Number)	(filing date)

I hereby claim the benefit under Title 35, United States Code, §120 of any United States application(s), or §365(c) of any PCT International application(s) designating the United States of America listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior United States or PCT International application in the manner provided by the first paragraph of Title 35, United States Code, §112, I acknowledge the duty to disclose material information as defined in Title 37, Code of Federal Regulations, §1.56



Attorney Docket No. C1043/7030

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the specification of which is attached hereto unless the following is checked:

KX was filed on January 12, 2001 as United States Application No. 09/743,621 or PCT
(Include Series Code)
International Application No. _____, bearing attorney docket No.
C1043/7030, and was amended on _____ (if applicable).

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(Number)	(Country-if PCT, so indicate)	(DD/MM/YY Filed)	YES NO
<u> </u>	<u> </u>	<u> </u>	<input type="checkbox"/> <input type="checkbox"/>
(Number)	(Country-if PCT, so indicate)	(DD/MM/YY Filed)	YES NO
<u> </u>	<u> </u>	<u> </u>	<input type="checkbox"/> <input type="checkbox"/>
(Number)	(Country-if PCT, so indicate)	(DD/MM/YY Filed)	YES NO

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<u> </u>	<u> </u>
(Application Number)	(filing date)
<u> </u>	<u> </u>
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which became available between the filing date of the prior application and the national or PCT International filing date of this application:

PCT/GB99/02271 (Application No.)	14 July 1999 (filing date)	Pending (status-patented, pending, abandoned:
(Application No.)	(filing date)	(status-patented, pending, abandoned:

PCT International Applications designating the United States:

(PCT Appl. No.)	(U.S. Ser. No.)	(PCT Filing date)	(status-patented, pending, abandoned)
-----------------	-----------------	-------------------	---------------------------------------

I hereby appoint the following attorney(s) and/or agent(s) to prosecute this application and to transact all business in the Patent and Trademark Office connected therewith:

David Wolf	17,528	Richard F. Giunta	36,149	William G. Gosz	27,787
George L. Greenfield	17,756	Douglas R. Wolf	36,971	Neil P. Ferraro	39,188
Stanley Sacks	19,900	Elizabeth R. Plummer	36,637	Lisa E. Winsor	44,405
Edward F. Perlman	28,105	Timothy J. Oyer	36,628	Mark Steinberg	40,829
Lawrence M. Green	29,384	John N. Anastasi	37,765	Stephen R. Finch	42,534
Steven J. Henry	27,900	Helen C. Lockhart	39,248	Joseph Teja Jr.	45,157
Edward R. Gates	31,616	James M. Hanifin, Jr.	37,929	Jeffrey B. Powers	45,021
Therese A. Hendricks	30,389	Christopher S. Schultz	39,039	Alan W. Steele	45,128
William R. McClellan	29,409	Paul D. Sorkin	40,212	Daniel P. McLoughlin	46,066
Ronald J. Krandsdorf	20,004	John R. VanAmsterdam	33,228	Robert H. Walat	46,324
M. Lawrence Oliverio	30,915	Matthew B. Lowrie	36,904	Thomas G. Field	45,596
Jason M. Honeyman	31,624	Robert E. Rigby, Jr.	41,316	Michael J. Pomianek	46,190
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Peter C. Lando	34,654	Robert M. Abrahamsen	37,482	Lisa K. Jorgenson	34,845
Gary S. Engelson	35,128	Ivan D. Zitkovsky	43,299	Robert D. McCutcheon	38,717
Peter J. Gordon	35,164	Alan B. Sherr	42,147	Jeffrey D. Moy	39,307
Randy J. Pritzker	35,986	Edward J. Russavage	43,069		

Address all correspondence to

Therese A. Hendricks
Wolf, Greenfield & Sacks, P.C.,
Federal Reserve Plaza
600 Atlantic Avenue
Boston, MA 02210-2211

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

Inventor's signature:	Date
Full name of sole or first inventor: Peter HO	
Citizenship: Singapore	
Residence: St. John's College, Cambridge CB2 1TP, United Kingdom	
Post Office Address: St. John's College, Cambridge CB2 1TP, United Kingdom	

12.4.07
Inventor's signature: _____


Date _____

Full name of second or joint inventor: Nir TESSLER

Citizenship: Israel

Residence: EE Dept., Technion, Haifa 32000, Israel

Post Office Address: ~~St. John's College, Cambridge, United Kingdom~~

As above

Inventor's signature: _____

Date _____

Full name of third joint inventor: Richard Henry FRIEND

Citizenship: British

Residence: 37 Barton Road, Cambridge CB3 9LG, United Kingdom

Post Office Address: 37 Barton Road, Cambridge CB3 9LG, United Kingdom

Inventor's signature: _____

Date _____

Full name of fourth joint inventor:

Citizenship:

Residence:

Post Office Address:

Inventor's signature: _____

Date _____

Full name of fifth joint inventor:

Citizenship:

Residence:

Post Office Address:

Inventor's signature: _____

Date _____

Full name of sixth joint inventor:

Citizenship:

Residence:

Post Office Address:



Attorney Docket No. C1043/7030

DECLARATION FOR PATENT APPLICATION

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I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled

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(Include Series Code)
International Application No. _____, bearing attorney docket No. C1043/7030, and was amended on _____ (if applicable).

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			[X] []
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(Number)	(Country-if PCT, so indicate)	(DD/MM/YY Filed)	
			[] []
<u> </u>	<u> </u>	<u> </u>	YES NO
(Number)	(Country-if PCT, so indicate)	(DD/MM/YY Filed)	
			[] []
<u> </u>	<u> </u>	<u> </u>	YES NO
(Number)	(Country-if PCT, so indicate)	(DD/MM/YY Filed)	

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<u> </u>	<u> </u>
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(Application No.)	(filing date)	(status-patented, pending, abandoned:

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Therese A. Hendricks	30,389	Christopher S. Schultz	39,039	Alan W. Steele	45,128
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Ronald J. Krandsdorf	20,004	John R. VanAmsterdam	33,228	Robert H. Walat	46,324
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James H. Morris	34,681	Robert A. Skrivaneck, Jr.	40,886	Theodore E. Galanthay	24,122
Peter C. Lando	34,654	Robert M. Abrahamsen	37,482	Lisa K. Jorgenson	34,845
Gary S. Engelson	35,128	Ivan D. Zitkovsky	43,299	Robert D. McCutcheon	38,717
Peter J. Gordon	35,164	Alan B. Sherr	42,147	Jeffrey D. Moy	39,307
Randy J. Pritzker	35,986	Edward J. Russavage	43,069		

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Citizenship: Singapore	
Residence: St. John's College, Cambridge CB2 1TP, United Kingdom	
Post Office Address: St. John's College, Cambridge CB2 1TP, United Kingdom	

Inventor's signature: _____
Full name of second or joint inventor: Nir TESSLER
Citizenship: Israel
Residence: EE Dept., Technion, Haifa 32000, Israel
Post Office Address: St. John's College, Cambridge, United Kingdom

Date

R.H. Friend

10 Feb 2001

Inventor's signature: _____
Full name of third joint inventor: Richard Henry FRIEND
Citizenship: British
Residence: 37 Barton Road, Cambridge CB3 9LG, United Kingdom
Post Office Address: 37 Barton Road, Cambridge CB3 9LG, United Kingdom

Date

Inventor's signature: _____
Full name of fourth joint inventor:
Citizenship:
Residence:
Post Office Address:

Date

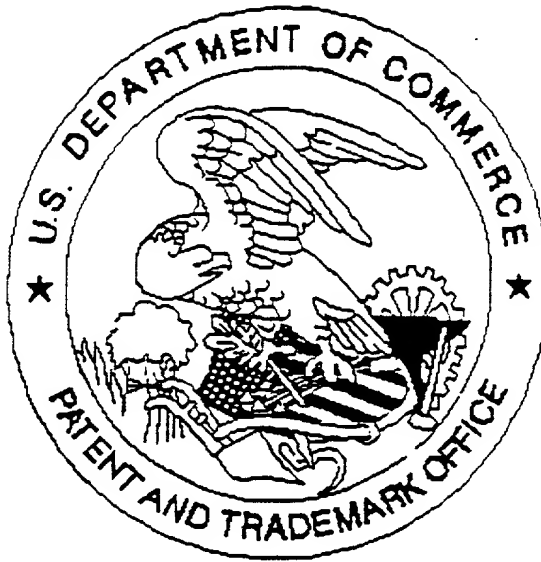
Inventor's signature: _____
Full name of fifth joint inventor:
Citizenship:
Residence:
Post Office Address:

Date

Inventor's signature: _____
Full name of sixth joint inventor:
Citizenship:
Residence:
Post Office Address:

Date

United States Patent & Trademark Office
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